



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) Publication number : **0 486 290 A2**

(12)

EUROPEAN PATENT APPLICATION

(21) Application number : **91310494.9**

(22) Date of filing : **14.11.91**

(51) Int. Cl.⁵ : **C12N 15/81, C12N 15/53,
C12P 5/02, C12P 33/00,
C12N 1/19, C12N 1/18,
// (C12N1/18, C12R1:865,
C12N1:19, C12R1:865)**

(30) Priority : **15.11.90 US 613380
28.10.91 US 783861**

(43) Date of publication of application :
20.05.92 Bulletin 92/21

(84) Designated Contracting States :
BE CH DE DK ES FR GB IT LI NL

(71) Applicant : **AMOCO CORPORATION
200 East Randolph Drive
Chicago Illinois 60601 (US)**

(72) Inventor : **Saunders, Court A.
210 Holmes
Claredon Hills, IL 60514 (US)
Inventor : Wolf, Fred Richard
912 Muirhead Avenue
Naperville, IL 60565 (US)
Inventor : Mukharji, Indrani
2300 Central Park Avenue
Evanston, IL 60201 (US)**

(74) Representative : **Horton, Sophie Emma et al
Elkington and Fife Prospect House 8
Pembroke Road
Sevenoaks, Kent TN13 1XR (GB)**

(54) **A method and composition for increasing the accumulation of squalene and specific sterols in yeast.**

(57) A method of increasing the accumulation of squalene and specific sterols in yeast comprising increasing the expression level of a structural gene encoding a polypeptide having HMG-CoA reductase activity in a mutant yeast having single or double defects in the expression of sterol biosynthetic enzymes is provided. The expression level of a structural gene is preferably increased by transforming yeast with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes a polypeptide having HMG-CoA reductase activity and a promoter that is suitable for driving the expression of the encoded polypeptide in the transformed yeast. The polypeptide having HMG-CoA reductase activity is preferably a truncated, active HMG-CoA reductase enzyme. Recombinant DNA molecules useful for transforming yeast and mutant yeast transformed with such recombinant DNA molecules are also disclosed.

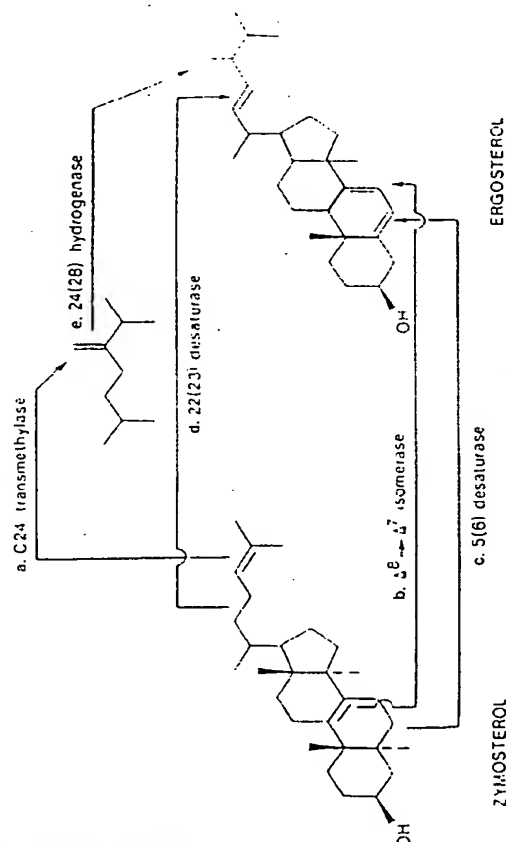


FIGURE 1

EP 0 486 290 A2

Jouve, 18, rue Saint-Denis, 75001 PARIS

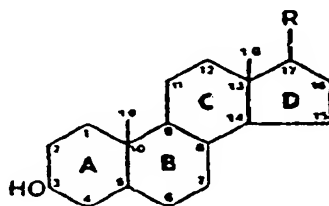
BEST AVAILABLE COPY

Technical Field

The present invention relates to a method and composition for increasing the accumulation of squalene and specific sterols in yeast. Squalene and sterol accumulation is increased by increasing the expression level of a gene encoding a polypeptide having HMG-CoA reductase activity.

Background of the Invention

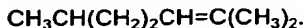
As used herein, the term "sterol" refers to derivatives of a fused, reduced ring system, cyclopenta-[α]-phenanthrene, comprising three fused cyclohexane rings (A, B and C) in a phenanthrene arrangement, and a terminal cyclopentane ring (D) having the formula and carbon atom position numbering shown below:



where R is an 8 to 10 carbon-atom sidechain.

Sterols are metabolically derived from acetate. Acetyl coenzyme A (CoA) reacts with acetoacetyl CoA to form 3-hydroxy-3-methylglutaryl CoA (HMG-CoA). HMG-CoA is reduced to mevalonate in an irreversible reaction catalyzed by the enzyme HMG-CoA reductase. Mevalonate is phosphorylated and decarboxylated to isopentenyl-pyrophosphate (IPP). Through the sequential steps of isomerization, condensation and dehydrogenation, IPP is converted to geranyl pyrophosphate (GPP). GPP combines with IPP to form farnesyl pyrophosphate (FPP), two molecules of which are reductively condensed to form squalene, a 30-carbon precursor of sterols.

In yeast, squalene is converted to squalene epoxide, which is then cyclized to form lanosterol. Lanosterol has two methyl groups at position 4, a methyl group at position 14, a double bond at position 8(9) and an 8 carbon sidechain at the formula:



Lanosterol is sequentially demethylated at positions 14 and 4 to form zymosterol (cholesta-8,24-dienol), which is converted to ergosterol (ergosta-5,7,22-trienol), the most abundant sterol of naturally occurring, wild-type yeast via a series of five enzymatic reactions schematically diagrammed in Figure 1.

The five reactions are:

- methylation of the carbon at position 24, catalyzed by a 24-methyltransferase;
- movement of the double bond at position 8(9) to position 7(8), catalyzed by a $\Delta 8-\Delta 7$ isomerase;
- introduction of a double bond at position 5(6), catalyzed by a 5-dehydrogenase (desaturase);
- introduction of a double bond at position 22(23), catalyzed by a 22-dehydrogenase (desaturase); and
- removal of a double bond at position 24(28), catalyzed by a 24(28)-hydrogenase (reductase).

In wild-type yeast of the species *Saccharomyces cerevisiae* (*S. cerevisiae*), the predominant order of these reactions is thought to be a, b, c, d and e. [Parks et al., *CRC Critical Reviews in Microbiology*, 6:301-341 (1978)].

According to such a predominant pathway, zymosterol is converted sequentially to fecosterol [ergosta-8,24(28)-dienol], episterol [ergosta-7,24(28)-dienol], ergosta-5,7,24(28)-trienol, ergosta-5,7,22, 24(28)-tetraenol, and finally ergosterol.

If the enzymes catalyzing the reactions involved in the predominant pathway are substrate specific, then one would expect to find only the six sterols set forth above in yeast. Such, however, is not the case. Eighteen sterols have been found and described. [See, e.g., Parks et al., *CRC Critical Reviews in Microbiology*, 6:301-341 (1978); Woods et al., *Microbios*, 10(A):73-80 (1974); Bard et al., *Lipids*, 12:645-654 (1977) (See Table 1)]. Thus, at least some of the enzymes are not substrate specific.

Table 1

	<u>Sterol</u>	<u>Required*</u> <u>Enzymes</u>
5	1. Zymosterol (cholesta-8,24-dienol)	none
10	2. fecosterol (ergosta-8,24(28)-dienol)	a
	3. episterol (ergosta-7,24(28)-dienol)	a,b
15	4. ergosta-5,7,24(28)-trienol	a,b,c
	5. ergosta-5,7,22,24(28)-tetraenol	a,b,c,d
20	6. ergosterol (ergosta-5,7,22-trienol)	a,b,c,d,e
25	7. ergosta-7,22,24(28)-trienol	a,b,d
	8. cholesta-7,24-dienol	b
30	9. cholesta-5,7,24-trienol	b,c
	10. cholesta-5,7,22,24-tetraenol	b,c,d
35	11. ergosta-5,7-dienol	a,b,c,e
	12. ergosta-7,22-dienol	a,b,d,e
40	13. ergosta-7-enol	a,b,e
	14. ergosta-5,8-dienol	a,c,e
	15. ergosta-5,8,22-trienol	a,c,d,e
45	16. ergosta-8,22-dienol	a,d,e
	17. ergosta-8-enol	a,e
50	18. ergosta-8,14,24(28)-trienol	a

* Enzymes theoretically required for the synthesis of the designated sterol.

Despite the lack of substrate specificity, one might expect that specific alterations in the sterol biosynthetic pathway would have predictable consequences. Currently available data show that such predictability is not

present.

For example, mutant *S. cerevisiae* with a defect in the expression of zymosterol-24-methyl-transferase (enzyme a), which mutants are designated erg6, might be expected to accumulate sterols 1 and 8-10 of Table 1, which sterols theoretically do not require the action of enzyme a for their synthesis. Parks et al., CRC Critical Reviews in Microbiology, 6:301-341 (1978), however, report that erg6 mutants accumulate only zymosterol (#1), cholesta-5,7,24-trienol (#9) and cholesta-5,7,22,24-tetraenol (#10). Bard, M. et al., Lipids, 12:645-654 (1977), on the other hand, report that erg6 mutants accumulate only sterols #1 and #10.

Mutant *S. cerevisiae* with a defect in the expression of ergosta-5,7,24(28)-trienol-22-dehydrogenase (enzyme d), designated erg5, might be expected to accumulate sterols 1-4, 6, 8, 9, 11, 13, 14, 17 and 18. Parks et al., CRC Critical Reviews in Microbiology, 6:301-341 (1978) report, that erg5 mutants accumulate only ergosta-5,7-dienol (#11), ergosta-5,7,24(28)-trienol (#4), ergosta-8,14,24(28)-trienol (#18) and episterol (#3). In contrast, Bard et al., Lipids, 12:645-654 (1977) report that erg5 mutants accumulate zymosterol (#1), ergosta-5,7-dienol (#11), ergosta-5,7,24(28)-trienol (#4), ergosta-7,24(28)-dienol (#3) and ergosta-8,14,24(28)-trienol (#18).

Still further, mutant *S. cerevisiae* with a defect in episterol-5-dehydrogenase (enzyme c), designated erg3, might be expected to accumulate sterols 1-3, 7, 8, 12, 13 and 16-18. Parks et al., CRC Critical Reviews in Microbiology, 6:301-341 (1978) report that erg3 mutants accumulate only ergosta-7,22-dienol (#12), ergosta-8,22-dienol (#16), ergosta-7,22,24(28)-trienol (#7), fecosterol (#2) and episterol (#3).

These data, taken together, show that specific defects in the expression of one sterol synthetic enzyme do not lead to predictable changes in sterol accumulation. A similar degree of unpredictability is found when sterol accumulation is examined in mutants having two defects in enzymes of the sterol biosynthetic pathway.

Thus, for example, erg5-erg6 double mutants (defects in enzymes d and a) might be expected to accumulate sterols 1, 8 and 9. Parks et al. and Bard et al., above, report that erg5-erg6 double mutants accumulate only zymosterol (#1) and cholesta-5,7,24-trienol (#9).

These data relating to sterol accumulation in yeast show that specific alterations in enzyme activity do not result in predictable changes in sterol accumulation. The data further show a lack of agreement between different investigators studying identical alterations. The present invention furnishes a solution to the problem of unpredictability by providing a method and composition for increasing the accumulation of squalene and specific sterols in yeast.

Summary of the Invention

The present invention generally provides a method of increasing squalene and specific sterol accumulation in mutant yeasts having a single or double defect in the expression of sterol biosynthetic pathway enzymes. This method comprises transforming such mutant yeasts with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes a polypeptide having HMG-CoA reductase activity and a promoter suitable for driving the expression of HMG-CoA reductase in the transformed yeast.

The structural gene encoding a polypeptide having HMG-CoA reductase activity preferably encodes an active, truncated HMG-CoA reductase enzyme, which enzyme comprises the catalytic and at least a portion of the linker region that is free from the membrane binding region of HMG-CoA reductase enzyme. The copy number of the structural gene is increased by transforming a mutant yeast with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes a polypeptide having a HMG-CoA reductase activity and a promoter that is suitable for driving the expression of the encoded polypeptide in the transformed yeast.

Suitable promoters include promoters that are subject to inducible regulation by factors either extrinsic or intrinsic to yeast. Preferably, both the promoter and the exogenous DNA segment are integrated into the chromosomal DNA of the transformed yeast.

The present invention most preferably provides a method of increasing squalene, zymosterol, cholesta-7,24-dienol and cholesta-5,7,24-trienol accumulation in yeast of the species *S. cerevisiae* comprising increasing the expression level of a structural gene encoding a polypeptide having HMG-CoA reductase activity in a mutant *S. cerevisiae* having defects in the expression of zymosterol-24-methyltransferase (erg6) and ergosta-5,7,24(28)-trienol-22-dehydrogenase (erg5).

In further preferred embodiments, transformation of a mutant yeast having a defect in the expression of the enzyme episterol-5-dehydrogenase (erg 3) results in a transformed, mutant yeast which overaccumulates squalene, ergosta-8,22-dienol, ergosta-7,22-dienol, ergosta-8-enol and ergosta-7-enol. Transformation of a mutant yeast having a double defect in the expression of zymosterol-24-methyltransferase and episterol-5-dehydrogenase enzymes (erg6 and erg3) results in a transformed mutant yeast which overaccumulates squalene, zymosterol and cholesta-7,24-dienol. Transformation of a mutant yeast having a defect in the exp-

ression of ergosta-5,7,24(28)-trienol-22-dehydrogenase (erg 5) results in a transformed mutant yeast which overaccumulates zymosterol and a mixture of ergosta-5,7,24(28)-trienol and ergosta-5,7-dienol.

Transformation of mutant yeast is preferably accomplished using a recombinant DNA molecule selected from the group of plasmid vectors consisting of plasmids pSOC725ARC, pSOC106ARC, pARC306E, pARC300D, pARC300S, pARC300T and pARC304S. Most preferred is plasmid pARC304S.

The present invention further provides for a mutant species of *S. cerevisiae*, which mutant has a double defect in the expression of zymosterol-24-methyl-transferase and ergosta-5,7,24(28)-trienol-22-dehydrogenase enzymes (erg 5 and erg6). That mutant *S. cerevisiae* is designated ATC0402mu.

The present invention still further provides for a mutant species of *S. cerevisiae* having a single or double defect in the expression of enzymes that catalyze the conversion of squalene to ergosterol that is transformed with a recombinant DNA molecule comprising as described before.

The present invention still further provides for recombinant DNA molecules used to transform mutant yeasts such that the transformed mutant yeast overaccumulates squalene and specific sterols. Preferred recombinant DNA molecules are plasmids pARC304S, pARC300S, pARC300T, pARC300D, pARC306E, pSOC106ARC and pSOC725ARC.

The present invention provides several benefits and advantages.

One advantage of the present invention is the provision of methods known to result in the predictable accumulation of specific sterols.

Another advantage of the present invention is the ability to accumulate specific sterols to levels markedly greater than levels found in non-transformed yeast.

Still further benefits and advantages will be apparent to the skilled worker from the description that follows.

Brief Descriptions of the Drawings

Figure 1 is a schematic diagram illustrating the various transformation steps involved in the metabolic conversion of zymosterol to ergosterol as shown and discussed in Bard et al., *Lipids*, 12(8):645 (1977). The letters (a-e) indicate the five enzymes responsible for catalyzing the individual transformation steps. Numerals alone or with the letter "C" and an enzymic name indicate the position of the enzymes' activities and the activity of each enzyme.

Figure 2, shown as twelve panels designated Figure 2-1 through 2-12, is the nucleotide base sequence (SEQ ID NO:1) and derived amino acid residue sequence (SEQ ID NO:2) for *S. cerevisiae* HMG-CoA reductase 1 published by Basson et al., *Mol. Cell Biol.* 8(9):3797-3808 (1988). Nucleotides are numbered (left-hand side) in the 5' to 3' direction. Position 1 corresponds to the first nucleotide of the ATG triplet coding for the initiator methionine. The predicted amino acid sequence is shown below the nucleotide sequence. The amino acid residues are numbered (right-hand side) beginning with the initiator methionine.

Figure 3 is a schematic diagram showing the physical structure and genetic organization of plasmid pSOC725ARC. Plasmid pSOC725ARC was constructed to place a coding sequence for a truncated HMG-CoA reductase gene under control of a GAL 1-10 promoter. This plasmid also contains the TRP-1 gene and the yeast 2 micron origin of replication. Certain restriction sites indicated by lines linked to the arcs and abbreviation for their respective restriction endonuclease enzymes are indicated.

Figure 4 is a schematic diagram showing the physical structure and genetic organization of plasmid pSOC106ARC. Plasmid pSOC106ARC was constructed to place a coding sequence for an intact HMG-CoA reductase gene under the control of a GAL 1-10 promoter. Plasmid pSOC106ARC also contains the TRP-1 gene and the yeast 2micron origin of replication. Certain restriction sites are indicated as in Figure 3.

Figure 5 is a schematic diagram showing the physical structure and genetic organization of plasmid pARC306E. Plasmid pARC306E was constructed to place a coding sequence for a truncated HMG-CoA reductase gene under control of a GAL-1 promoter. Plasmid pARC306E also contains the TRP-1 gene. Certain restriction sites are indicated as in Figure 3.

Figure 6 is schematic diagram showing the physical structure and genetic organization of plasmid pARC300D. Plasmid pARC300D was constructed to place a coding sequence for a truncated HMG-CoA reductase gene under the control of a PGK promoter. Plasmid pARC300D also contains the TRP-1 gene. Certain restriction sites are indicated as in Figure 3.

Figure 7 is a schematic diagram showing the physical structure and genetic organization of plasmid pARC300S. Plasmid pARC300S was constructed to place a coding sequence for a truncated HMG-coA reductase gene under control of a PGK promoter. Plasmid pARC300S also contains a URA 3 selectable marker. Certain restriction sites are indicated as in Figure 3.

Figure 8 is a schematic diagram showing the physical structure and genetic organization of plasmid pARC300T. Plasmid pARC300T was constructed to place a coding sequence for a truncated HMG-coA reduc-

tase gene under control of a PGK promoter. Plasmid pARC300T also contains a URA 3 selectable marker. Certain restriction sites are indicated as in Figure 3.

Figure 9 is a schematic diagram showing the physical structure and genetic organization of plasmid pARC304S. Plasmid pARC304S was constructed to place a coding sequence of a truncated HMG-CoA reductase gene under the control of an ADH promoter. Plasmid pARC304S also contains a URA 3 selectable marker. Certain restriction sites are indicated as in Figure 3.

Detailed Description of the Invention

I. Definitions

The following words and phrases have the meanings set forth below.

Expression: The combination of intracellular processes, including transcription and translation, undergone by a structural gene to produce a polypeptide.

Expression vector: A DNA sequence that forms control elements that regulate expression of structural genes when operatively linked to those genes.

Operatively linked: A structural gene is covalently bonded in correct reading frame to another DNA (or RNA as appropriate) segment, such as to an expression vector so that the structural gene is under the control of the expression vector.

Promoter: A recognition site on a DNA sequence or group of DNA sequences that provide an expression control element for a structural gene and to which RNA polymerase specifically binds and initiates RNA synthesis (transcription) of that gene.

Recombinant DNA molecule: A hybrid DNA sequence comprising at least two nucleotide sequences not normally found together in nature.

Structural gene: A DNA sequence that is expressed as a polypeptide, i.e., an amino acid residue sequence.

Vector: A DNA molecule capable of replication in a cell and/or to which another DNA segment can be operatively linked so as to bring about replication of the attached segment. Alternatively, a vector can be a non-replicating vector that is integrated into the chromosome of the transformed cell. A plasmid is an exemplary vector.

II. The Invention

The present invention relates to compositions and methods for increasing the accumulation of squalene and specific sterols in yeast cultures as well as to the yeast that exhibit increased squalene and sterol accumulation relative to a non-transformed yeast. Preferred yeasts are yeasts of the Saccharomyces or Candida genus. A more preferred yeast is Saccharomyces cerevisiae (S. cerevisiae).

A yeast contemplated by this invention is transformed with an added structural gene that encodes a polypeptide having HMG-CoA reductase activity, that encoded polypeptide being expressed in the transformed yeast. Preferred non-transformed yeasts are mutant species having a single or double defect in the expression of enzymes involved in converting zymosterol to ergosterol (sterol biosynthetic pathway enzymes). The non-transformed and transformed yeasts compared are of the same species, such as S. cerevisiae.

Sterol production in a yeast culture of the present invention is increased by increasing the cellular activity of the enzyme HMG-CoA reductase, which enzyme catalyzes the conversion of 3-hydroxy-3-methylglutaryl Coenzyme A (HMG-CoA) to mevalonate. As used herein, "cellular activity" means the total catalytic activity of HMG-CoA reductase in a yeast cell.

Cellular HMG-CoA reductase activity is increased by increasing the expression level of a structural gene encoding a polypeptide having HMG-CoA reductase catalytic activity. Expression of that encoded structural gene enhances the cellular activity of that enzyme. The expression level is increased by methods well known in the art. For example, expression of a structural gene is increased by deregulating the promoter, which controls expression of such a structural gene. The promoter that regulates expression of the HMG-CoA reductase gene in a normal, wild-type yeast can be identified and excised from the genome. A new promoter, which allows for overexpression of the HMG-CoA reductase gene, is then inserted according to standard transformation techniques. A preferred means of increasing the expression level of a structural gene encoding a polypeptide having HMG-CoA reductase catalytic activity is to increase the copy number of a structural gene encoding such a polypeptide.

The copy number is increased by transforming a yeast cell with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes a polypeptide having HMG-CoA reduc-

tase activity, and a promoter suitable for driving the expression of said polypeptide in said yeast. Such a polypeptide is catalytically active, and is preferably a truncated HMG-CoA reductase protein.

Thus, a transformed yeast cell has one or more added genes that encode a polypeptide having HMG-CoA reductase activity relative to a non-transformed yeast of the same species. As such, a transformed yeast can be distinguished from a non-transformed yeast by standard technology such as agarose separation of DNA fragments or mRNAs followed by transfer and appropriate blotting with DNA or RNA or by use of polymerase chain reaction technology, as are well known. Relative HMG-CoA reductase activity of the transformed and non-transformed yeasts can also be compared, with a relative increase in HMG-CoA reductase activity in transformed yeasts being indicative of transformation.

The accumulation of squalene and specific sterols can also be used to distinguish between non-transformed and transformed yeasts.

A. Structural Genes

The present invention contemplates transforming a yeast with a structural gene that encodes a polypeptide having HMG-CoA reductase activity. The HMG-CoA reductase enzymes of both animal and yeast cells comprise three distinct amino acid residue sequence regions, which regions are designated the catalytic region, the membrane binding region and the linker region.

The catalytic region contains the active site of the HMG-CoA reductase enzyme and comprises about forty percent of the total, localized on the COOH-terminal portion of intact HMG-CoA reductase enzyme. The membrane binding region contains hydrophobic amino acid residues and comprises about fifty percent of the total, localized on the NH₂-terminal portion of intact HMG-CoA reductase enzyme. The linker region connects the catalytic and membrane binding regions, and constitutes the remaining about ten percent of the intact enzyme.

As discussed in greater detail below, only the catalytic region of HMG-CoA reductase is needed herein. Thus, a structural gene that encodes a polypeptide corresponding to that catalytic region is the minimal gene required for transforming yeasts. However, larger polypeptide enzymes and their structural genes are preferred. Thus, the present invention contemplates use of truncated structural genes that encode the active catalytic region, or the catalytic region plus at least a portion of the linker region that is free from the membrane binding region of HMG-CoA reductase.

A structural gene encoding a polypeptide having HMG-CoA reductase activity can be obtained or constructed from a variety of sources and by a variety of methodologies, [See, e.g., Carlson et al., *Cell*, 28:145 (1982); Rine et al., *Proc. Nat. Acad. Sci. U.S.A.*, 80:6750 (1983)]. Exemplary of such structural genes are the mammalian and yeast genes encoding HMG-CoA reductase.

The mammalian genome contains a single gene encoding HMG-CoA reductase. The nucleotide base sequence of the hamster and human gene for HMG-CoA reductase have been described. A composite nucleotide sequence of cDNA corresponding to the mRNA, as well as the derived amino acid residue sequence, for hamster HMG-CoA reductase is found in Chin et al., *Nature*, 308:613 (1984) and SEQ ID NO:3. The composite nucleotide sequence in that paper, comprising about 4606 base pairs, includes the nucleotide sequence encoding the intact hamster HMG-CoA reductase enzyme.

Intact hamster HMG-CoA reductase comprises about 887 amino acid residues, shown in SEQ ID NO:4.

A preferred structural gene is one that encodes a polypeptide corresponding to only the catalytic region of the enzyme. Two catalytically active segments of hamster HMG-CoA reductase have been defined, [Liscum et al., *J. Biol. Chem.*, 260(1):522 (1985)]. One catalytic region has an apparent size of about 63 kDa and comprises amino acid residues from about position 373 to about position 887 of SEQ ID NO:4. A second catalytic region has an apparent size of about 53 kDa and comprises amino acid residues from about position 460 to about position 887 of SEQ ID NO:4. The about 63 kDa catalytically active segment is encoded by base pairs from about nucleotide position 1282 to about nucleotide position 2824 of the sequence in SEQ ID NO:3. The about 53 kDa catalytically active segment is encoded by base pairs from about nucleotide position 1543 to about nucleotide position 2824 of the sequence in SEQ ID NO:3.

In a preferred embodiment, the utilized structural gene encodes the catalytic region and at least a portion of the linker region of HMG-CoA reductase. The linker region of hamster HMG-CoA reductase comprises amino acid residues from about position 340 to about position 373 or from about position 340 to about position 460, depending upon how the catalytic region is defined. These linker regions are encoded by base pairs from about nucleotide position 1183 to about nucleotide position 1282 or from about position 1183 to about position 1543 respectively of the sequence in SEQ ID NO:3. The structural gene encoding the linker region is operatively linked to the structural gene encoding the catalytic region.

In one particularly preferred embodiment, a structural gene encoding a catalytically active, truncated HMG-CoA reductase enzyme can optionally contain base pairs encoding a small portion of the membrane region of

the enzyme. A truncated hamster HMG-CoA reductase gene, designated HMGR- Δ 227, comprising nucleotides 164-190 and 1187-2824 of the sequence in SEQ ID NO:3, which encodes amino acid residues 1-9 (from the membrane binding region) and 342-887 has been used to transform cells lacking HMG-CoA reductase, [Gil et al., *Cell*, 41:249 (1985)].

5 A structural gene encoding a polypeptide comprising a catalytically active, truncated or intact HMG-CoA reductase enzyme from other organisms such as yeast can also be used in accordance with the present invention.

Yeast cells contain two genes encoding HMG-CoA reductase. The two yeast genes, designated HMG1 and HMG2, encode two distinct forms of HMG-CoA reductase, designated HMG-CoA reductase 1 and HMG-CoA reductase 2. The nucleotide base sequence of HMG1 (SEQ ID NO:1) as well as the amino acid residue sequence of HMG-CoA reductase 1 (SEQ ID NO:2) are presented in Figure 2, reprinted from Basson et al., *Mol. Cell Biol.*, 8(9):3797 (1988).

The entire HMG1 gene comprises about 3360 base pairs. Intact HMG-CoA reductase 1 comprises an amino acid sequence of about 1054 amino acid residues.

15 The entire HMG2 gene comprises about 3348 base pairs shown in SEQ ID NO:5. Intact HMG-CoA reductase 2 comprises about 1045 amino acid residues shown in SEQ ID NO:6 (Basson et al., above).

By analogy to the truncated hamster structural gene, structural genes encoding polypeptides comprising catalytically active, truncated HMG-CoA reductase enzymes from yeast can also be used in accordance with the present invention.

20 The catalytic region of HMG-CoA reductase 1 comprises amino acid residues from about residue 618 to about residue 1054: i.e., the COOH-terminus. A structural gene that encodes the catalytic region comprises base pairs from about nucleotide position 1974 to about position 3282 of Figure 2 and SEQ ID NO:1.

The linker region of HMG-CoA reductase 1 comprises an amino acid sequence from about residue 525 to about residue 617. A structural gene that encodes the linker region comprises nucleotides from about position 1695 to about position 1974 of Figure 2. A structural gene encoding a polypeptide comprising the catalytic region and at least a portion of the linker region of yeast HMG-CoA reductase 1 preferably comprises the structural gene encoding the linker region of the enzyme operatively linked to the structural gene encoding the catalytic region of the enzyme.

25 Also by analogy to the truncated hamster gene, a truncated HMG1 gene can optionally contain nucleotide base pair sequences encoding a small portion of the membrane binding region of the enzyme. Such a structural gene preferably comprises base pairs from about nucleotide position 121 to about position 146 and from about position 1695 to about position 3282 of Figure 2 and SEQ ID NO:1.

A construct similar to those above from an analogous portion of yeast HMG-CoA reductase 2 can also be utilized.

35 It will be apparent to those of skill in the art that the nucleic acid sequences set forth herein, either explicitly, as in the case of the sequences set forth above, or implicitly with respect to nucleic acid sequences generally known and not presented herein, can be modified due to the built-in redundancy of the genetic code and non-critical areas of the polypeptide that are subject to modification and alteration. In this regard, the present invention contemplates allelic variants of structural genes encoding a polypeptide having HMG-CoA reductase activity.

40 The previously described DNA segments are noted as having a minimal length, as well as total overall lengths. That minimal length defines the length of a DNA segment having a sequence that encodes a particular polypeptide having HMG-CoA reductase activity. As is well known in the art, so long as the required DNA sequence is present and in proper reading frame, (including start and stop signals), additional base pairs can be present at either end of the segment and that segment can still be utilized to express the protein. This, of course, presumes the absence in the segment of an operatively linked DNA sequence that represses expression, expresses a further product that consumes the enzyme desired to be expressed, expresses a product other than the desired enzyme or otherwise interferes with the structural gene of the DNA segment.

45 Thus, so long as the DNA segment is free of such interfering DNA sequences, the maximum size of a recombinant DNA molecule, particularly an expression vector, is governed mostly by convenience and the vector size that can be accommodated by a host cell, once all of the minimal DNA sequences required for replication and expression, when desired, are present. Typically, a DNA segment of the invention can be up to 15,000 base pairs in length. Minimal vector sizes are well known.

55 B. Recombinant DNA Molecules

A recombinant DNA molecule of the present invention can be produced by operatively linking a vector to a useful DNA segment to form a plasmid such as discussed herein. Particularly preferred recombinant DNA

molecules are discussed in detail in Examples 2 to 7, hereafter. A vector capable of directing the expression of a polypeptide having HMG-CoA reductase activity is referred to herein as an "expression vector".

Such expression vectors contain expression control elements including the promoter. The polypeptide coding genes are operatively linked to the expression vector to permit the promoter sequence to direct RNA polymerase binding and expression of the desired polypeptide coding gene. Useful in expressing the polypeptide coding gene are promoters that are inducible, viral, synthetic, constitutive as described by Poszkowski et al., *EMBO J.*, 3:2719 (1989) and Odell et al., *Nature*, 313:810 (1985), and temporally regulated, spatially regulated, and spatiotemporally regulated as disclosed in Chau et al., *Science*, 244:174-181 (1989). The promoter preferably comprises a promoter sequence whose function in regulating expression of the structural gene is substantially unaffected by the amount of sterol in the cell. As used herein, the term "substantially unaffected" means that the promoter is not responsive to direct feedback control by the sterols accumulated in transformed cells.

A promoter is also selected for its ability to direct the transformed yeast's transcriptional activity to the structural gene encoding a polypeptide having HMG-CoA reductase activity. Structural genes can be driven by a variety of promoters in yeast.

Promoters utilized with the present invention are those preferably regulated by factors, which can be monitored and controlled in the internal or external environment of the transformed cell. Examples of promoters inducibly regulated by factors in the cell's external environment (extrinsic factors) are the GAL 1 promoter, the GAL 10 promoter, the GAL 1-10 promoter, the GAL 7 promoter, the metallothionein promoter, the α -factor promoter, the invertase promoter and the enolase promoter. Preferred are the well known GAL 1, the GAL 10 and the GAL 1-10 promoters.

Examples of promoters subject to inducible regulation by factors in the cell's internal environment (intrinsic factors) are the phosphoglycerate kinase (PGK) promoter, the triose-phosphate isomerase (TPI) promoter, the alcohol dehydrogenase (ADH) promoter and the repressible acid phosphatase promoter. Preferred are the well known PGK and the ADH promoters.

The choice of which expression vector and ultimately to which promoter a polypeptide coding gene is operatively linked depends directly on the functional properties desired, e.g. the location and timing of protein expression, and the host cell to be transformed. These are well known limitations inherent in the art of constructing recombinant DNA molecules. However, a vector useful in practicing the present invention is capable of directing the expression of the polypeptide coding gene included in the DNA segment to which it is operatively linked.

The present method contemplates a plasmid vector. The plasmid vectors of the present invention can be incorporated either within (integrated) or without (episomal) the chromosomes of the transformed cell. An episomal plasmid includes an origin of replication for yeast, the nucleic acid sequence that encodes a polypeptide having HMG-CoA reductase activity, a promoter, and a selective marker. The selective marker can include genes conveying antibiotic resistance, or permitting an auxotrophic host to metabolize a substrate that it would not otherwise be able, but for the presence of the plasmid vector. However, the use of antibiotic resistance as a selective marker requires growing organisms in an antibiotic culture media. Due to the expense of the antibiotic, organisms dependent on antibiotics are difficult to develop commercially. Generally, auxotrophic organisms are used for yeast.

Auxotrophic organisms can be produced by mutation and culture techniques which are well known in the art. Selective markers which can complement an auxotrophic host organism include the well known TRP 1 gene encoding phosphoribosyl anthraniline isomerase, the URA 3 gene encoding orotidine-5' phosphate decarboxylase, the LEU 2 gene encoding isopropylmalate isomerase, and the HIS 3 gene encoding histidinol dehydrogenase. A preferred selective marker for an auxotrophic host is TRP 1. Preferred episomal plasmid vectors are pSOC725ARC and pSOC106ARC.

Episomally replicating vectors are sometimes difficult to maintain in host organisms for long periods of time in liquid culture, especially when the selective pressure used to maintain the vector is complementation of a nutritional auxotrophy. A preferred embodiment of the present invention includes an integrating vector which requires little or no selective pressure to maintain base sequences for the polypeptide having HMG-CoA reductase activity and the promoter.

Integrating vectors, in accordance with the present invention, include base sequences that encode a polypeptide having HMG-CoA reductase activity, a promoter, a selective marker and sequences homologous to host chromosomal DNA that permit the base sequences to be incorporated within the chromosome via homologous recombination. The homologous region includes restriction sites that permit the plasmid to become linear. In linear form, the plasmid can recombine at homologous regions of the chromosome. Integrating vectors do not include origins of replication for the host organism.

Preferred integrating vectors are pARC300S, pARC300T, pARC300D, pARC306E and pARC304S. Plasmid vector pARC304S is most preferred as evidenced by its ability to generate the greatest enhancement in

sterol accumulation (see Example 15). The basic genetic characteristics of preferred plasmid vectors are summarized in Table 2, below.

TABLE 2

<u>Plasmid Vector</u>	<u>Genetic Characteristics</u>
pSOC106	TRP1-2 μ ori-GAL 1-HMG1*
pSOC725	TRP1-2 μ ori-GAL 10-tHMG1**
pARC306E	TRP1-GAL 1-tHMG1
pARC300D	TRP1-PGK-tHMG1
pARC300S,T	URA3-PGK-tHMG1-ura3 term
pARC304S	URA3-ADH-tHMG1-ura3 term

* HMG1 - gene encoding intact S. cerevisiae HMG-CoA reductase 1.

** tHMG1 - gene encoding catalytic region and a portion of the linker region of S. cerevisiae HMG-CoA reductase 1.

Individuals skilled in the art will readily recognize that episomal and integrating vectors are often amplified in organisms other than the intended host and require means of replication and selection in the non-host organism. Generally, the non-host organism is Escherichia coli due to its well-known features and characteristics.

In preferred embodiments, the vector used to express the polypeptide coding gene includes a selection marker that is effective in a yeast cell, such as the URA 3 or TRP 1 markers. Other suitable selection means for use in amplifying the vectors in bacteria include antibiotic markers, such as genes encoding for beta lactamase (penicillin resistance), chloramphenicol transacetylase (chloramphenicol resistance), and neomycin phosphotransferase (kanamycin and neomycin resistance).

A variety of methods has been developed to operatively link DNA to vectors via complementary cohesive termini or blunt ends. For instance, complementary homopolymer tracts can be added to the DNA segment to be inserted and to the vector DNA. The vector and DNA segment are then joined by hydrogen bonding between the complementary homopolymeric tails to form recombinant DNA molecules.

Alternatively, synthetic linkers containing one or more restriction endonuclease sites can be used to join the DNA segment to the expression vector. The synthetic linkers are attached to blunt-ended DNA segments by incubating the blunt-ended DNA segments with a large excess of synthetic linker molecules in the presence of an enzyme that is able to catalyze the ligation of blunt-ended DNA molecules, such as bacteriophage T4 DNA ligase. Thus, the products of the reaction are DNA segments carrying synthetic linker sequences at their ends. These DNA segments are then cleaved with the appropriate restriction endonuclease and ligated into an expression vector that has been cleaved with an enzyme that produces termini compatible with those of the synthetic linker. Synthetic linkers containing a variety of restriction endonuclease sites are commercially available from a number of sources including New England BioLabs, Beverly, MA.

Also contemplated by the present invention are RNA equivalents of the above described recombinant DNA molecules.

C. Transformed Yeasts and Methods of Transformation

The copy number of a gene coding for a polypeptide having HMG-CoA reductase activity is increased by transforming a desired yeast with a suitable vector that contains that structural gene. Expression of that gene in the transformed yeast enhances the activity of HMG-CoA reductase.

Yeast cells are transformed in accordance with the present invention by methods known and readily apparent to those of skill in the yeast transformation art, [See, e.g., Hinnen et al., Proc. Natl. Acad. Sci. USA, **75**:1929-1978; Ito et al., Bact., **5**:163-168 (1983)].

A preferred general method of transformation is the lithium acetate procedure of Ito et al., above. Yeast cells are grown to a concentration about 2×10^7 cells/ml in a medium containing yeast extract, bactopectone and dextrose. Cells are collected by low speed centrifugation and resuspended in a transformation medium containing lithium acetate in a Tris-EDTA buffer.

Cells are maintained in the transformation medium for about one hour at about 30°C. Recombinant DNA molecules of desired composition are added to the transformation medium cell suspension and the mixture is maintained at about 30°C for about one-half hour. Polyethylene glycol (M.W. 4000) is then added to the cell suspension such that the final concentration of polyethylene glycol is about 35 percent weight/volume (w/v). Cells are maintained in the polyethylene glycol-containing solution at about 30°C for about two hours and then at about 42°C for an additional five minutes. Sterile distilled water is added to the cell suspension, and the cells collected by low speed centrifugation. Further specifics are provided hereinafter.

Successfully transformed cells are identified by growing the transformed cells on selection medium, identifying cell characteristics indicative of transformation (i.e., increased accumulation of squalene or specific sterols), analyzing nucleic acids isolated from such transformed cells with standard techniques such as Southern blot analysis, [Holm et al., Gene, **42**:169 (1986)].

D. Mutated Yeasts

The yeasts utilized in accordance with the present invention are mutated yeasts having single or double defects in the expression of enzymes that catalyze the conversion of zymosterol to ergosterol. Such enzymes are referred to herein as "erg" gene products. Table 3 below lists the particular erg designations for specific enzyme expression defects.

Table 3

<u>Enzyme Expression Defect</u>	<u>Mutant Designation</u>
zymosterol-24-methyltransferase	erg6
ergosta-5,7,24(28)-trienol-22-dehydrogenase	erg5
episterol-5-dehydrogenase	erg3

Mutants used in accordance with the present invention can be purchased or generated from commercially available sources such as the Yeast Genetic Stock Center (Berkeley, CA.). For example, erg5 and erg5-erg6 double mutants are produced from commercially available sources.

Mutant yeast ATC0402mu, an erg5-erg6 double mutant, is constructed by crossing a commercially available erg6 mutant yeast, M610-12B, with a commercially available erg5 mutant, pol5 Δ 22, and then crossing the resultant double mutant, ATC0403mu, with a wild-type yeast. Mutant yeast ATC0402mu and its derivative mutant yeast ATC0315rc are the most preferred mutants for transformation with the plasmid vectors of the present invention.

Alternatively, ATC0403 is crossed with a different wild-type, and mutants having desired genotypes are back-crossed twice with wild-type yeast to yield species ATC4124, an erg5 mutant.

Mutants are also obtained by well known methods of inducing mutations. See, e.g., Boeke et al., Mol. Gen. Genet., **197**:345-346 (1984); Sherman et al., Methods and Yeast Genetics, Cold Spring Harbor Laboratory, N.Y. (1986).

In a preferred embodiment, wild-type yeasts are transformed with an inducible "TY1-neo" transposon as a mutagenic agent. Plasmid pJEF1105, containing a GAL:TY1-neo expression cassette, is used as the transforming agent. Boeke et al., Science **239**:280-282 (1989). Competent transformants demonstrating both neomycin and nystatin resistance are then evaluated for sterol content.

Transformation of wild-type yeast with pJEF1105 yields mutant ATC6118, an erg3 mutant, and mutant ATC0501, an erg6 mutant.

Mutants having single expression defects are then crossed to generate mutants having double defects in enzyme expression. For example, the crossing of mutant ATC6118 with mutant ATC0501 yields mutant

ATC6119, an erg3-erg6 double mutant.

The genotype of exemplary mutants contemplated for use in the present invention are presented in Table 4 below. Genotype symbols are used in accordance with convention cited in Mortimer et al. *Yeast*, 5:321-403 (1989) and Broach, *The Molecular Biology of the Yeast Saccharomyces, Life Cycle and Inheritance*, Strathern, Jones and Broach, eds., Cold Spring Harbor Laboratory, pp. 653-727 (1981).

Table 4

<u>Species</u>	<u>genotype</u>
pol5 α Δ 22	a, erg5
M610-12 β	α , ile3, erg6-5, trp1, gal2
DBY745	α , adel, ura3-52, leu2-100, leu2-122,
	MEL, gal 1 gal 10
YNN281	α , trp1- Δ , his3 Δ -200, ura 3-52, lys 2
ATC0403mu	a, trp1, gal, erg5, erg6
ATC0402mu	a, trp1, GAL, erg5, erg6
ATC6118	a, his3 Δ -200, erg3, ura3-52, GAL
ATC4124	α , erg5, trp1, GAL
ATC4154	a, ura3-52, erg7, gal
ATC6119	α , erg3, erg6, ura3-52, GAL
ATC1500cp	a, erg5, erg6
ATC0315rc	a, ura3, erg5, erg6
ATC1551	a, erg5, erg6

E. Squalene and Sterol Accumulation in Transformed Yeast

The transformed mutant yeast species of the present invention overaccumulate squalene and specific sterols relative to non-transformed mutants of the same species. Relative to a non-transformed erg3 mutant, an erg3 mutant transformed with a plasmid vector used herein overaccumulates squalene, ergosta-8,22-dienol, ergosta-7,22-dienol, ergosta-8-enol and ergosta-7-enol.

Relative to a non-transformed erg5 mutant, an erg5 mutant transformed with a plasmid vector used herein overaccumulates squalene, zymosterol, and a mixture of ergosta-5,7,24(28)-trienol and ergosta-5,7 dienol.

Similar results are seen when mutants having double defects in enzymes of the sterol synthetic pathway are transformed. Relative to a non-transformed erg3-erg6 mutant, an erg3-erg6 mutant transformed with a useful plasmid vector overaccumulates squalene, zymosterol and cholesta-7,24-dienol.

Relative to a non-transformed erg5-erg6 mutant, an erg5-erg6 double mutant transformed with the plasmid vector useful herein overaccumulates squalene, zymosterol, cholesta-5,7,24-trienol and cholesta-7,24-dienol.

F. HMG-CoA Reductase Activity In Transformed Yeasts

The expression of a structural gene encoding a polypeptide having HMG-CoA reductase activity in the transformed yeast of the present invention enhances the cellular activity of said HMG-CoA reductase. As a result of transformation, the copy number of an added gene encoding a polypeptide having HMG-CoA reductase activity is increased from 1 to about 2 to about 10.

Cellular activity of HMG-CoA reductase in such transformed cells is almost linearly proportional to the increase in copy number through a copy number of about 6 and then falls slightly when a copy number of 9 is reached. Thus, when the copy number is increased to about 2, HMG-CoA reductase activity is elevated to a level about 1.4 times the activity observed in non-transformed yeast. A further increase in the copy number to a level of about 6 is accompanied by a further increase in HMG-CoA reductase activity to a level about 2.6 times that found in non-transformed yeast. Increases in the copy number beyond about 6 to about 9 are not accompanied by further increases in HMG-CoA reductase activity. A transformed yeast having a copy number

of about 9 has a level of HMG-CoA reductase activity about equal to about twice that seen in non-transformed yeast.

G. Harvesling of Sterols

If desired, transformed yeasts are harvested to recover the sterol product. Most of the sterol in our genetically transformed yeast of this invention occurs in the form of fatty acid esters. To obtain free sterols, it is therefore necessary to saponify the "yeast pulp" in base, e.g., as described in the Examples below (2:1 EtOH/H₂O containing 20 percent w/v KOH).

In a preferred embodiment, harvesting comprises:

- (i) homogenizing sterol-containing transformed yeasts to produce a pulp; and
- (ii) extracting the sterol(s) from the pulp with an appropriate basic solvent such as an organic solvent or by supercritical extraction followed by base saponification in an appropriate solvent [Favati et al., *J. Food Sci.*, 53:1532 (1988) and the citations therein] to produce a sterol-containing liquid solution or suspension; and
- (iii) isolating the sterol(s) from the solution or suspension.

Transformed yeasts are homogenized to produce a pulp using methods well known to one skilled in the art. This homogenization can be done manually, by a machine, or by a chemical means. The pulp consists of a mixture of the sterol of interest, residual amounts of precursors, cellular particles and cytosol contents, which is subjected to extraction procedures.

Sterol(s) can be extracted from the pulp produced above to form a sterol-containing solution or suspension. Such extraction processes are common and well known to one skilled in this art. For example, the extracting step can consist of soaking or immersing the pulp in a suitable solvent. This suitable solvent is capable of dissolving or suspending the sterol present in the pulp to produce a sterol-containing solution or suspension. Solvents useful for such an extraction process are well known to those skilled in the art and include several organic solvents and combinations thereof such as methanol, ethanol, isopropanol, acetone, acetonitrile, tetrahydrofuran (THF), hexane, and chloroform as well as water-organic solvent mixtures. A vegetable oil such as peanut, corn, soybean and similar oils can also be used for this extraction.

Yeasts transformed with a structural gene for an active, truncated HMG-CoA reductase enzyme are grown under suitable culture conditions for a period of time sufficient for sterols to be synthesized. The sterol-containing yeast cells are then lysed chemically or mechanically, and the sterol is extracted from the lysed cells using a liquid organic solvent, as described before, to form a sterol-containing liquid solution or suspension. The sterol is thereafter isolated from the liquid solution or suspension by usual means such as chromatography.

The sterol is isolated from the solution or suspension produced above using methods that are well known to those skilled in the art of sterol isolation. These methods include, but are not limited to, purification procedures based on solubility in various liquid media, chromatographic techniques such as column chromatography and the like.

Best Mode For Carrying Out The invention

The following examples illustrate the best mode of carrying out the invention and are not to be construed as limiting of the specification and claims in any way.

EXAMPLE 1: Transformation of *S. Cerevisiae*

Yeast of the species *S. cerevisiae* were transformed in accordance with a lithium acetate procedure, [Ito et al., *J. Bacteriol.*, 153:163-168 (1983)]. Yeast cells were grown in about 50 ml of YEPD medium (yeast extract 1 percent w/v, bacto-peptone, 2 percent w/v; and dextrose, 2 percent w/v) overnight at about 30°C. When the concentration of cells was about 2×10^7 cells/ml, the cells were collected by low speed centrifugation. Cells appearing in the pellet of the centrifugation were suspended in about 50 mls of TE buffer (10 mM Tris-Cl, 1 mM EDTA) and repelleted by centrifugation. The pellet from this second centrifugation was resuspended in about 1.0 ml of TE buffer. To 0.5 ml of this cell suspension were added 0.5 ml of 0.2 M lithium acetate (LiOAc), and the suspension was maintained at about 30°C for one hour with constant shaking.

Recombinant DNA (about 10 µg in up to 15 µl of TE buffer) was added to 100 µl of the TE-LiOAc cell suspension and the admixture maintained at about 30°C for one-half hour without shaking. The DNA-containing cell suspension was then well mixed with polyethylene glycol (44 percent w/v) such that the final concentration of polyethylene glycol (PEG) was about 35 percent (w/v).

The cells were maintained in this PEG solution at about 30°C for about two hours and then at about 42°C

for about five minutes. About 10 ml of sterile, distilled water was added to each suspension and the cells were collected by low speed centrifugation. This procedure was repeated, and the collected cells were dispersed in about 1.0 ml of distilled water. Approximately 100 to 200 µl of this suspension were then spread-plated on selective medium.

- 5 Transformation of cells was confirmed by growth on selection medium, identification of cell characteristics indicative of transformation (i.e., increased levels of selected sterols or squalene), and Southern blot analysis of nucleic acid isolated from such transformed cells [Holm et al., Gene, 42:169-173 (1986)].

EXAMPLE 2: Construction of Episomal Plasmid pSOC725ARC

10

Plasmid pSOC725ARC (See Figure 3) was constructed to place a coding sequence for a truncated HMG1 gene under control of the GAL 1 portion of a GAL 1-10 promoter. Plasmid pSOC725ARC also contains the TRP 1 gene and the yeast 2 micron origin of replication (IR1). This plasmid was prepared from intermediate plasmids as follows.

- 15 The TRP 1-ARS gene of S. cerevisiae was removed from plasmid YRP12 [Stinchcomb et al. Nature, 282:39 (1979)] by digestion with Eco RI. The 1445 base pair DNA fragment containing the TRP 1-ARS gene was purified on an agarose gel and ligated into plasmid pUC8 (Viera et al., Gene, (1982)), which had been digested with Eco RI to form plasmid pSOC742.

- 20 A yeast episomal replication origin, obtained from purified S. cerevisiae two-micron plasmid DNA, was digested with Eco RI and then treated with the Klenow fragment of E. coli DNA polymerase 1 to yield an about 2240 base pair fragment containing the two-micron origin of DNA replication. The about 2240 base pair fragment was purified by agarose gel electrophoresis and ligated into plasmid pUC8, which had been digested with Sma I to form plasmid pSOC743.

- 25 Plasmid pSOC742 was cleaved with Bam HI and Bgl II to yield an 857 base pair, TRP 1-containing gene fragment, which was inserted into pSOC743 that had been cut with Bam HI to form plasmid pSOC744.

The MEL1 gene was removed from plasmid pMP550 [Summer-Smith et al., Gene, 36:333-340 (1985)] with restriction endonucleases Eco RI and Bam HI, and the about 2858 base pair restriction fragment containing MEL1 was purified on an agarose gel. The purified fragment was then ligated into plasmid pUC8 which had been digested with Eco RI and Bam HI to form plasmid pSOC741.

- 30 The final stage of assembly of pSOC740 was achieved by purifying an about 3101 base pair, Eco RI restriction fragment of pSOC744 that contained the TRP 1 and two-micron origin, and ligating it into Eco RI-cleaved plasmid pSOC741 to form plasmid pSOC740.

- 35 The GAL 1-10 promoter was excised from pBM258, [Johnston et al., Proc. Natl. Acad. Sci. USA, 79:6971-6975 (1982)] as a 685 base pair Bam HI-Eco RI restriction fragment, and ligated into pUC18, which had been digested with Bam HI and Eco RI to form plasmid pSOC711.

Plasmid pSOC740 was digested with Eco RI and the resulting 3101 base pair fragment, containing the two-micron origin of replication and the TRP 1 gene, was isolated and ligated into the Eco RI digested plasmid pSOC711 to produce plasmid pSOC712, in which the TRP 1 gene is proximal to the GAL 1-10 promoter.

- 40 A Pst I restriction site spanning the coding sequence for amino acid residues 529-530 of HMG-coA reductase 1 was chosen as the point at which to introduce both a new Bam HI restriction site and a new initiator methionine codon. A 1706 base pair Pst I-Eco RI restriction fragment, containing the coding sequence for the COOH-terminal half of HMG-CoA reductase 1, was purified from a digest of pJR59, [Basson et al., Proc. Natl. Acad. Sci. USA, 83:5563-5567 (1986)]. This purified pJR59 fragment and a synthetic oligonucleotide:

- 45 **d5' -GATCCGTCGACGCATGCCTGCA-3' (SEQ ID NO:7)**
d3' -GCAGCTGCGTACGG-5' (SEQ ID NO:8)

- 50 were ligated with pUC18 [Yanisch-Perron et al., Gene, 33:103-119 (1985)], which had been cleaved with Bam HI and Eco RI.

The resulting plasmid, pSOC937, contained a Bam HI restriction site 12 base pairs upstream of the truncated HMG-CoA reductase coding sequence initiator methionine. The polypeptide formed from initiation at that point had initial methionine and proline residues followed by amino acid residues 530 through 1054 of the natural HMG-CoA reductase 1.

- 55 The Eco RI restriction site, which is at the 3' end of the gene, is located 135 base pairs past the end of the coding sequence for the truncated HMG-CoA reductase protein. The truncated gene for HMG-CoA reductase was placed into plasmid pSOC712 by converting the Eco RI site at the 3' end of the truncated reductase gene to a Bam HI site (Klenow polymerase filled, ligated to an oligonucleotide, d5-CGGATCCG, specifying the Bam

HI restriction site) and cleaving the preparation with endonuclease Bam HI. A purified, resulting 1728 base pair Bam HI ended restriction fragment from pSOC937 was ligated into the Bam HI-digested pSOC712 to produce plasmid pSOC725ARC, whose schematic restriction map is shown in Figure 3.

5 EXAMPLE 3: Construction of Episomal Plasmid pSOC106ARC

Plasmid pSOC106ARC (See Figure 4) was constructed to place a coding sequence for intact HMG1 under the control of the GAL 1 portion of a GAL 1-10 promoter.

10 A 610 base pair Bgl II fragment from pJR59 (about positions 9026-9636), containing the DNA surrounding the beginning of the HMG-CoA reductase coding sequence, was isolated and further restricted with Dde I to provide a DNA fragment (about positions 9151-9636) starting 68 base pairs upstream of the first codon of the HMG-CoA reductase coding sequence.

15 The Dde I and Bgl II fragments were treated with the Klenow fragment of DNA polymerase to render the ends "blunt." The fragments were then ligated to oligonucleotide linkers, d5'-CCGGATCCGG-3' (SEQ ID NO:9), specifying a Bam HI cleavage site (BRL linkers). The ligated fragments were digested with Bam HI to produce ligatable Bam HI restriction ends, and the resulting 499 base pair fragment containing the start of the HMG-CoA reductase coding sequence was ligated into Bam HI digested pBR322 to form plasmid pSOC104.

20 The remainder of the HMG-CoA reductase coding sequence was reconstructed downstream of the new 5' Bam HI site by ligating a 1477 base pair Xba I-Sac I DNA fragment of pJR59, which specifies the 5' half of the HMG-CoA reductase coding sequence, and a 2101 base pair Sac I-Sal I fragment of pJR59, which specifies the 3' half of the HMG-CoA reductase coding sequence, into pSOC104 digested with Xba I and Sal I to form plasmid pSOC105 containing a 3903 base pair Bam HI-Sal I restriction fragment having the entire coding sequence for HMG-CoA reductase. This 3903 base pair fragment was ligated into Bam HI-Sal I-restricted pSOC712 (See Example 2) to form plasmid pSOC106ARC.

25 EXAMPLE 4: Construction of Integrating Plasmid pARC306E

Plasmid pARC306E (See Figure 5) was constructed to place a coding sequence for truncated HMG1 under control of the GAL 1 portion of a GAL 1-10 promoter.

30 Plasmid pARC306E contains the S. cerevisiae TRP 1 gene and a GAL 1 promoter-driven, truncated HMG-CoA reductase gene housed on an E. coli replicon, which specifies ampicillin resistance. There are no S. cerevisiae replicators on plasmid pARC306E. Unique restriction sites within both the TRP 1 gene (Eco RV, position 865) and the truncated HMG-CoA reductase gene (Cla I, position 4280) serve as sites for the generation of linear plasmids with DNA homologous to S. cerevisiae chromosomal DNA on both sides of the restriction site. Thus, plasmid pARC306E can be incorporated into the chromosome at either site via homologous recombination.

The multiple restriction recognition site of plasmid pUC8, located between the Eco RI and Hind III sites, was replaced by the oligonucleotide:

40 d5' -AGCTTTTCGCGAGCTCGAGATCTAGATATCGATG (SEQ ID NO:10)
3' -AGCGCTCGAGCTCTAGATCTATAGCTACTTAA-5'
(SEQ ID NO:11)

45 to create plasmid pUC8NL, which has a single restriction site for the nuclease enzyme Cla I.

Plasmid pSOC712 (See Example 2) was digested with Eco RI and the fragments treated with nuclease S1 and bacteriophage T4 DNA polymerase plus deoxynucleotides to remove the overhanging 5' Eco RI restriction ends. These ends were ligated to the oligonucleotide:

d5'-CATCGATG-3'
d3'-GTAGCTAC-5'

50 and the fragments treated with Cla I nuclease to produce Cla I restriction ends.

The resulting 3108 base pair Cla I-Cla I fragment, containing the yeast TRP 1 gene and the two-micron replicator, was purified by gel electrophoresis and ligated into pUC8NL, which had been cleaved with Cla I, to create plasmid pARC300A.

55 A 2031 base pair fragment containing the two-micron replication origin was removed from pARC300A by treatment with nuclease Pst I. The resulting modified plasmid pARC300A was treated with nuclease S1 and bacteriophage T4 DNA polymerase plus deoxynucleotides to remove the Pst I restriction overhangs and with calf intestinal alkaline phosphatase to disallow reclosure of the plasmid. The modified pARC300A plasmid was

coligated with the oligonucleotide:

d5'-CATCGATG-3'
d3'-GTAGCATC-5'

to introduce a Cla I site just downstream (to the 3' end) of the TRP 1 gene to form a plasmid, and then closed to form pARC306B. The TRP 1 gene was separate from yeast replicators, and bounded by Cla I restriction sites.

Plasmid pARC306B was digested with Cla I, purified by polyacrylamide gel electrophoresis and the Cla I-Cla I restriction fragment was introduced into plasmid pUC8, which had been cleaved with nuclease Acc I, to form plasmid pARC306C.

As the integration of exogenous DNA into yeast chromosomes is best carried out using homologous recombination, a dispensable fragment of yeast DNA was desired. This DNA would be used to drive homologous recombination if for some reason, recombination at the TRP 1 or HMG-CoA reductase gene were not utilizable. The DNA chosen for this purpose was the HIS3 gene.

An 1800 pair Bam HI-Bam HI restriction fragment was removed from plasmid YEP6 [Struhl et al., *Proc. Natl. Acad. Sci. USA*, 76:1035 (1979)] and introduced into plasmid pARC306C, which had been cleaved with Bam HI, to create plasmid pARC306D. Plasmid pSOC725 (See Example 2) was digested with Eco RI to yield a GAL 1-10 promoter linked to a truncated HMG-CoA reductase gene, which was then inserted into Eco RI-digested plasmid pARC306D, to form plasmid pARC306E.

EXAMPLE 5: Construction of Integrating Plasmid PARC300D

Plasmid pARC300D (See Figure 6) was constructed to place a coding sequence for a truncated HMG1 gene under the control of a PGK promoter. This plasmid was prepared from intermediate plasmids as follows.

Plasmid pSOC611 was constructed to determine the efficacy of the mouse metallothionine promoter as a transcriptional driver for the truncated HMG-CoA reductase gene in yeast. Construction of pSOC611 began with restriction of plasmid pSOC744 (See Example 2) with Eco RI endonuclease, followed by treatment with Klenow Polymerase I and deoxynucleotide triphosphates to fill in the Eco RI restriction ends. The resulting about 3101 base pair 2-micron- and TRP 1-containing fragment of pSOC744 was ligated to pUC18 which had been cleaved with Hinc II, to form plasmid pSOC517.

Plasmid pSOC517 was then cleaved with Kpn I and Eco RI and the mouse metallothionine promoter was inserted as a Kpn I-Eco RI restriction fragment to form plasmid pSOC518. This promoter region is composed of the Kpn I to Bgl II fragment originally in pJYMMT (e) [Hammer et al., *Journal of Applied Molecular Genetics*, Vol. 1:273 (1982)] as well as a short Bgl II, Eco RI DNA fragment of unknown sequence.

The truncated HMG-CoA reductase gene was added to pSOC518 in two steps. First, the truncated HMG-CoA reductase gene was removed from pSOC725 as a Bam HI restriction fragment. This fragment was then ligated into M13mp7 which had been cleaved with Bam HI. The new M13 derivative formed was designated pSOC610. The truncated HMG-CoA reductase gene was removed from pSOC610 as an Eco RI fragment and inserted into Eco RI-digested plasmid pSOC518. The resulting plasmid was designated pSOC611.

Plasmid pUC8 was partially digested with restriction endonuclease Hae II and religated. Transformants arising from this procedure were screened to find a plasmid missing the Hae II restriction fragment containing the portion of the lac operon which was originally present in plasmid pUC8. This new plasmid was designated pSOC505ARC. Restriction sites for the endonucleases Eco RI, Hind III and Kpn I were introduced into the Nde I site of plasmid pSOC505ARC by ligation of the oligonucleotide:

d5'-TATCGAATTCAAGCTTGGTACCGA-3' (SEQ ID NO:12)

3'-AGCTTAAGTTCGAACCATGGCTAT-5' (SEQ ID NO:13)

into Nde I-digested pSOC505ARC to form plasmid pARC303A.

To form the new multi-cloning site, the normal multi-cloning site present in M13mp18 was altered by ligating the oligonucleotide:

d5'-GATCCAGCTGTGTAC-3' (SEQ ID NO:14)

d3'-GTCGACA-5'

into Bam HI-Kpn I digested M13mp18. This resulted in an altered M13 virus, designated pARC303B. This construct lacked both the Kpn I and Sma I sites normally found in the M13mp18 multi-cloning site. The new multi-cloning site was removed as an Eco RI, Hind III restriction fragment from pARC303B, and was ligated into

Eco RI, Hind III restricted plasmid pARC303A to form plasmid pARC303C.

In addition to a variation in the normal array of sites included in the multi-cloning site, another smaller multi-cloning site was introduced into the vector, at a point some distance away from the first multi-cloning site to allow for independent manipulation of yeast auxotrophic complementation markers and other features which did not have to be proximal to the promoters and coding sequences which would be inserted in the large multi-cloning site. The new array of restriction sites was introduced by ligation of the oligonucleotide:

d5'-CCCGGGATCGATCACGT-3' (SEQ ID NO:15)

d3'-TGCAGGGCCCTAGCTAG-5' (SEQ ID NO:16)

into pARC303C cleaved with endonuclease Aat II to form plasmid pARC300E, which contained the series of cloning sites, Aat II, Sma I, and Cla I at the former Aat II site.

The yeast TRP 1 gene was isolated as an 820 base pair fragment from pARC306B (See Example 4) with the restriction endonuclease Cla I. The 820 base pair Cla I-Cla I fragment was purified by agarose gel electrophoresis and ligated into plasmid pARC300E, which had been digested with Cla I, to create plasmid pARC300B.

Plasmid pSOC611 was digested with Bam HI and Ssp I to yield a 1667 base pair coding sequence for the truncated HMG-CoA reductase gene which was purified by agarose gel purification. The 1667 base pair fragment was ligated to Bam HI, Hinc II restricted plasmid pARC300B to generate plasmid pARC300C.

A source of an alternate promoter to the GAL 1-10 promoter which has been used to drive transcription of the truncated HMG-CoA reductase gene was desired. Use of the GAL 1-10 promoter requires that the yeast be cultured on galactose, an expensive substrate. In order to achieve high levels of transcription through the truncated HMG-CoA reductase gene during culture, growth in the presence of the much less expensive substrate, glucose, the promoter from the S. cerevisiae phosphoglycerate kinase (PGK) gene was isolated. The sequence of the gene is available from the literature, [Hitzeman, et al., Nucl. Acid Res., 10:7791-7808 (1982)].

From the known sequence, an oligonucleotide probe sufficiently complementary to the gene to be used as a hybridization probe was synthesized:

d5'-ATAAAGACATTGTTTTTTAGATCTGTTGTAA-3' (SEQ ID NO:17)

This probe was labelled by T₄ polynucleotide kinase treatment in the presence of ³²P-ATP, and used to screen a library of bacteriophage λ subclones of the yeast genome, supplied by Maynard Olson (Washington University School of Medicine, Department of Genetics, St. Louis, Mo.). The gene was removed from this clone as an Eco RI-Hind III fragment, and subcloned into M13mp18, forming a new phage mARC127.

To make the PGK promoter useful, the restriction site at the 5' end of the promoter was changed to an Eco RI restriction site, and a Bgl II restriction site was introduced into the DNA fragment to the 3' side of the transcriptional start site. The Bgl II restriction site was introduced by using the oligonucleotide:

d5'-ATAAAGACATTGTTTTTTAGATCTGTTGTAA-3' (SEQ ID NO:17),

to mutagenize mARC127 according to the procedure of Kunkel et al., Proc. Natl. Acad. Sci. USA, 82:4778 (1985). This resulted in the M13 phage designated mARC128.

The Hind III site beyond the 5' end of the promoter region was converted to an Eco RI site by cutting mARC128 with nuclease Hind III, treating with the Klenow fragment of DNA polymerase and the four deoxynucleotide triphosphates, followed by ligation in the presence of the oligonucleotide:

d5'-GGAATTCC-3',

which specifies an Eco RI site. The resulting M13 derivative was designated pARC306L.

Plasmid pARC306L was digested with Eco RI and Bgl II and a 1500 base pair fragment containing the PGK promoter, was purified by agarose gel electrophoresis and ligated into pARC300C, which had been restricted with Eco RI and Bam HI, to produce plasmid pARC300D.

EXAMPLE 6: Construction of Integrating Plasmids pARC300S and pARC300T

Plasmids pARC300S (See Figure 7) and pARC300T (See Figure 8) were constructed to incorporate a URA 3 selectable marker into an integrating vector, in which a coding sequence for a truncated HMG1 gene was under the control of a PGK promoter.

The only difference between plasmid pARC300S and pARC300T is the length of the PGK promoter driving

transcription of the truncated reductase coding sequence. A unique Eco RV restriction site found within the URA 3 gene allows the plasmids to be linearized and integrated via homologous recombination into the chromosomal URA 3 gene.

The URA 3 gene from plasmid YEP24 (Botstein, et al., Gene, 8:17-24 (1979)) was removed as an 1127 base pair Eco RI-Sma I ended restriction fragment and ligated into plasmid pUC19, cut with Eco RI and Sma I to form a new plasmid LpARCLH550. An 1141 base pair Hind III ended restriction fragment was removed from LpARCLH550 and ligated into Hind III-cleaved pUC18 to form plasmid LpARCLH553a. An 1108 base pair Sma I-Hind III restriction fragment was removed from LpARCLH553a and inserted into Sma I-Hind III cleaved M13mp19 nucleic acid to create a new phage nucleic acid pARC306K. The unique PstI site within the URA 3, gene was eliminated by mutagenesis with the oligonucleotide:

d5' GATTTATCTTCGTTTCCTGCAAGTTTTTGTTC-3' (SEQ ID NO:18),

using the method of Kunkel, L.M.; et al., Proc. Nat' l. Acad. Sci. USA, 82:4778 (1985), to form plasmid pARC300Z.

Plasmid pARC300Z was cut with Hind III, the ends filled in with the Klenow fragment of DNA polymerase and deoxynucleotide triphosphates, and the modified pARC300Z ligated with oligonucleotide d5'-CCCCGGGG-3', which specified a Sma I restriction site. This new M13 derivative, which contains the URA 3 gene on a Sma I restriction fragment, was named plasmid pARC300Y.

Plasmid pARC304A was constructed to provide a source of a modified URA 3 transcription terminator fragment which could then be introduced at the 3' end of the coding sequence region in a yeast integrating transformation vector. The transcription terminator would function to improve mRNA stability in species transformed with integrating vectors containing coding sequences either lacking the terminator or having only weak terminator sequences. Improved mRNA stability could mean increased activity of the protein encoded by the coding sequence region. The terminator chosen was a region of the S. cerevisiae URA 3, which functions as a terminator, [Yarger et al., Molecular and Cellular Biology, 6:1095 (1986)]. The terminator sequence was constructed using 4 synthetic oligomers:

d5' -AGCTTCGAAGAACGAAGGAAGGAGCACAGACTTAG-3' (SEQ ID NO:19)

d5' -ATTGGTATATATACGCATATTGCGGCCGCGGTAC-3' (SEQ ID NO:20)

d5' -CGCGGCCGCAATATGCGTATATATAC-3' (SEQ ID NO:21)

d5' -CAATCTAAGTCTGTGCTCCTTCCTTCGTTCTTCGA-3' (SEQ ID NO:22)

These oligomers were designed to provide Hind III and Kpn I restriction ends, respectively. The modified URA 3 transcription terminator was assembled by ligating all four oligomers to each other and digesting the ligation product with Hind III and Kpn I to produce ligatable Hind III-Kpn I restriction ends. The 67 base pair fragment was isolated on a polyacrylamide gel, purified by electroeluting the DNA from the gel fragment, and then ligated into Hind III-Kpn I restricted pUC118, (ATCC 37462). This construction created a new plasmid designated pARC304A.

A 67 base pair Hind III-Kpn I fragment containing a URA 3 transcription terminator was isolated from plasmid pARC304A and ligated into Hind III-Kpn I restricted pARC300E to form plasmid pARC300M. A truncated HMG-CoA reductase coding sequence was isolated as a 1667 base pair Bam HI-Ssp I fragment from pSOC611, (See Example 5) purified by agarose gel electrophoresis, and ligated to pARC300M, which had been restricted with Bam HI and Hinc II, to form plasmid pARC300R.

A URA 3 complementing gene was removed from plasmid pARC300Y as an Xma I restriction fragment, and ligated into the Xma I site of pARC300R to create plasmid pARC300U.

One other change in the restriction sites available on the DNA specifying the PGK promoter was made. The minimum DNA required to specify full PGK promoter activity has been determined, [Stanway, Nucleic Acids Research, 15:6855-6873 (1987)]. A new Eco RI site was added to the DNA specifying the PGK promoter at a

region just past the minimal 5' required DNA. The site was added by utilizing the oligonucleotide:

d5'-CTTTATGAGGGTAACATGAATTCAAGAAGG-3' (SEQ ID NO:23),

to mutagenize mARC1228 by the method of Kunkel et al., Proc. Natl. Acad. Sci. USA, 82:4778 (1985). This new M13 derivative was designated pARC306M.

A 1500 base pair phosphoglycerate kinase promoter (PGK) was removed from plasmid pARC306L (See Example 5) using Eco RI and Bgl II restriction enzymes. The PGK promoter fragment was purified by agarose gel electrophoresis and ligated to Eco RI and Bam HI restricted pARC300U, to form plasmid pARC300S.

A shortened PGK promoter (555 base pair fragment) was isolated from Eco RI and Bgl II restricted plasmid pARC306M and inserted into Eco RI-Bam HI digested plasmid pARC300U to form plasmid pARC300T.

The only difference between plasmid pARC300S and plasmid pARC300T is the length of the PGK promoter driving transcription of the truncated reductase coding sequence. A unique Eco RV restriction site found within the URA 3 gene allows the plasmids to be linearized and integrated via homologous recombination into the chromosomal URA 3 gene.

EXAMPLE 7: Construction of Plasmid pARC304S

Plasmid pARC304S (see Figure 9) was constructed to place the coding sequence of a truncated HMG1 gene under the control of an ADH promoter.

Plasmid pBR322 was digested with Eco RI and Bam HI to yield a fragment containing the ADH1 promoter. The ADH1-containing fragment was ligated into plasmid pARC300U (See Example 6), which had been cut with Eco RI and Bam HI, to form pARC304S.

Plasmid pARC304S was deposited pursuant to the Budapest Treaty requirements with the American Type Culture Collection (ATCC) at 12301 Parklawn Drive, Rockville, MD 20852 U.S.A. on November 9, 1990 and was assigned Accession No. ATCC40916.

EXAMPLE 8: Generation of Mutant S. cerevisiae ATC0402mu

Mutant ATC0402mu was generated to have the GAL, a, and trp1 phenotype as well as having defects in the expression of zymosterol-24-methyltransferase and ergosta-5,7,24(28)-trienol-22-dehydrogenase enzymes. These enzymes are respectively the erg6 and erg5 gene products of S. cerevisiae.

An erg6 deficient mutant S. cerevisiae, M610-12B, obtained from the Yeast Genetic Stock Center (Univ. of California, Berkeley, CA), was crossed with an erg5 deficient mutant S. cerevisiae (obtained as a gift from Dr. Leo Parks, North Carolina State Univ., Raleigh, NC) to produce an erg6-erg5 double mutant, ATC0403mu.

ATC0403mu was then crossed with wild-type S. cerevisiae, DBY745 (Yeast Genetic Stock Center) to produce mutant ATC0402mu.

Mutant ATC0402mu was deposited pursuant to the Budapest Treaty Requirements with the American Type Culture Collection (ATCC) at 12301 Parklawn Drive, Rockville MD 20852 U.S.A. on November 9, 1990, and was assigned Accession No. ATCC 74027.

EXAMPLE 9: Generation of Transformed Mutants ATC1500cp, ATC1502, ATC1503, ATC1551 and ATC2401

Several mutants were generated from the transformation of ATC0402mu using the method of Example 1, with various expression systems (plasmids) containing HMG-CoA reductase coding sequences under the transcriptional control of various promoters. The introduction into ATC0402mu of plasmid pSOC106ARC, constructed according to the method of Example 3, created ATC1503.

The introduction into ATC0402mu of plasmid pSOC725ARC, constructed according to the method of Example 2, created ATC2401mu.

The introduction into ATC0402mu of plasmid pARC306E, constructed according to the method of Example 4, created ATC1502.

The introduction into ATC0402mu of plasmid pARC300D, constructed according to the method of Example 5, created ATC1500cp.

The creation of strain ATC1551 required the generation of a ura3 derivative of strain ATC1500cp, which has no auxotrophic markers. The ura3 derivative was created by transforming ATC1500cp with a mutagenic oligonucleotide using the method of Moerschell et al. [Proc. Natl. Acad. Sci. USA, 85:524-528 (1988)]. The sequence of the mutagenic oligonucleotide used is:

5' -GCCAAGTAGTTTTTACTCTTCAAGACAGATAATTTGCTGACA-3'
(SEQ ID NO: 24)

5 Mutated yeast cells were selected by their resistance to 5'-fluoro-orotic acid (5-FOA), as described in Ausubel et al., (eds.), Current Protocols in Molecular Biology, John Wiley and Sons, New York, (1989), and screened for their inability to grow in the absence of uracil. The resulting ura3 strain was designated ATC0135rc. Strain ATC0315rc was then transformed with plasmid pARC304S, constructed according to the method of Example 7, to create strain ATC1551.

10 Transformation of strain ATC0315rc with plasmid pARC304S of the present invention resulted in the greatest degree of sterol accumulation. Further, the growth of a transformed ATC0315rc mutant under conditions of restricted aeration as compared to usual culture conditions, resulted in an increased accumulation of squalene relative to other sterols as well as an increase in the total accumulation of squalene and total sterols.

15 Mutant ATC0315rc was deposited pursuant to the Budapest Treaty Requirements with the American Type Culture Collection (ATCC) at 12301 Parklawn Drive, Rockville, MD 20852 U.S.A. on September 16, 1991, and was assigned Accession No. ATCC 74090.

EXAMPLE 10: Generation of Mutant *S. cerevisiae* ATC6118, ATC0501 and ATC6119

20 Mutants were obtained using an inducible "TY1-neo" transposon as the mutagenic agent, [Boeke, et al., Science, 239:280-282 (1989)].

Wild type *S. cerevisiae* JB516 was transformed with plasmid pJEF1105 [Boeke et al., Science, 239:280-282 (1989)], containing an inducible GAL:TY1neo expression cassette, and plasmid pCGS286, containing a GAL-lacZ control. The transformed yeast were then spread onto petri dishes containing two kinds of Xgal chromogenic indicator dye: synthetic dextrose (SD) agar media minus uracil and synthetic galactose (SG) agar media minus uracil. Yeast transformed with plasmid pJEF1105 appeared normal on dextrose but smaller than untransformed control yeast on galactose media.

The stability of plasmid pJEF1105 was confirmed by shuttling into *E. coli* for propagation and restriction analysis.

30 Once plasmid pJEF1105-transformed yeasts were shown to be competent, the pJEF1105 transformants were placed on SG-minus uracil agar at a density of no more than 1000 transformants per petri plate. The plates were incubated at 22°C for five days, during which the mutagenic transposition of the plasmid borne TY1-neo occurred. The transformants were then replica plated onto another SG-minus uracil plate and incubated another five days. Those colonies that survived were replica plated onto YEPD agar plates containing 100 units/ml of nystatin to select for sterol production and 100 units/ml of G418 (a neomycin analog) to select for the "neo" phenotype. Transformants that were both nystatin and G418 resistant were evaluated for sterol content and distribution using gas chromatographic and mass spectrographic analysis and then classified as to the specific sterol biosynthetic step affected by the mutation.

40 A yeast deficient in the enzyme episterol-5-dehydrogenase (the *erg3* gene product) was isolated and designated ATC6118.

A yeast deficient in the enzyme zymosterol-24-methyltransferase (*erg6*) was isolated from plasmid pJEF1105 mutated yeast DBY745 (Yeast Genetic Stock Center) and designated ATC0501.

ATC0501 was crossed with ATC6118 to produce an *erg3-erg6* double mutant designated ATC6119.

45 EXAMPLE 11: Generation of Transformed Mutant *S. cerevisiae* ATC2100, ATC2104 and ATC2109

Following the method of Example 1, the introduction into ATC6119 of plasmids pARC300S and pARC300T, constructed according to the method of Example 6, created ATC2100 and ATC2104 respectively, whereas the introduction into ATC6118 of plasmid pARC300S created ATC2109.

50 EXAMPLE 12: Generation of Mutant *S. cerevisiae* ATC4124

ATC4124 (Yeast Genetic Stock Centers) was generated by crossing ATC0403mu with YNN281 (Yeast Genetic Stock Centers) and selecting for the desired mutation. The resulting segregants were then backcrossed twice with YNN281.

55 Resulting ATC4124 had a defect in the expression of cholesta-5,7,24(28)-trienol-22-dehydrogenase (the *erg5* gene product).

EXAMPLE 13: Generation of Transformed Mutant *S. cerevisiae* ATC2107 and ATCE2108

Following the method of Example 1, introduction into ATC4124 of plasmid pARC306E, constructed according to the method of Example 4, created ATC2107 and ATC2108.

EXAMPLE 14: HMG-CoA Reductase Activity in Mutant and Transformed Yeast

HMG-CoA reductase activity was measured in non-transformed and transformed *erg5-erg6* mutant yeasts. About 0.2 ml of 50 mM potassium phosphate buffer, pH 6.8, containing 125 mM sucrose, 20 mM EDTA and 100 mM KCl was combined with 10 mM DTT (freshly made), 1 mM NADPH, enzyme preparation and water to make an enzyme solution of about 0.475 ml final volume. The enzyme solution was preincubated at 37°C for 20 minutes and the incubation reaction initiated with the addition of 100 μ M 14 C-HMG-CoA (60,000 dpm in 0.025 ml). After five minutes, the reaction was stopped by the addition of 50 μ l of HCl (1:1) and further incubation at 37°C for 30 minutes to lactonize the product. The product, mevalonolactone, was separated from HMG on an anion exchanger AGI-X8 (Bio-Rad) and the radioactivity associated with the product was counted in a scintillation counter. The results are shown in Table 5, below. The copy number of an added structural gene encoding a polypeptide having HMG-CoA reductase activity was estimated using standard procedures well known to those of skill in the transformation art.

TABLE 5

<u>Mutant</u>	<u>Estimated Copy # of Added Structural Gene</u>	<u>Specific Activity HMG-CoA Reductase (mmols/min/mg dry wt)</u>
<u>Non-transformed</u>		
ATC0402mu	0	0.52
<u>Transformed</u>		
ATC1503	1, 2	0.69
ATC1500cp	5, 6	1.33
ATC1512	8, 9	1.01

EXAMPLE 15: Squalene and Sterol Accumulation in Yeast

The accumulation of squalene and specific sterols was determined in non-transformed and transformed mutant yeast cultures.

Fifty to one hundred mg of lyophilized yeast cells were extracted/saponified in 10 ml of an ethanol/water (2:1) solution containing 20 percent (w/v) KOH for two hours at 80°C. Extracts were partially neutralized with 10 ml 1N HCl and extracted twice with 15 ml n-heptane. The sterol-containing heptane fractions were evaporated to dryness under a stream of N₂ and resuspended to an appropriate volume with n-heptane containing an internal standard (5- α -cholestane).

The resuspended samples were analyzed for sterol accumulation by capillary GC with flame ionization detection.

Table 6 contains summary data for non-transformed (control) and transformed mutants having a single defect (*erg3*, *erg5*) in the expression of sterol biosynthetic pathway enzymes.

Table 7 contains summary data for non-transformed (control) and transformed mutants having double defects (*erg3-erg6*, *erg5-erg6*) in the expression of sterol biosynthetic pathway enzymes.

In both Table 6 and Table 7, the transformants were all made by transforming the control mutant having the same *erg* mutation.

Sterol levels are expressed as a percent of the dry biomass.

TABLE 6
ERG3 Mutants

	<u>Sterol</u>	<u>Percent of Biomass</u>	
		<u>Non-transformed</u>	<u>Transformed</u>
		ATC6118	ATC2109
5	a. Squalene	N.D.*	0.26
10	b. ergosta-8,22-dienol	0.31	1.08
	c. ergosta-7,22-dienol	0.66	1.64
	d. ergosta-8-enol	0.27	0.42
15	e. ergosta-7-enol	0.63	0.72

ERG5 Mutants

	<u>Sterol</u>	<u>Percent of Biomass</u>	
		<u>Non-transformed</u>	<u>Transformed</u>
		ATC4124	ATC2107 ATC2108
20	a. Squalene	N.D.	1.10 0.49
	b. Zymosterol	0.05	0.25 0.25
25	c. ergosta-5,7, 24(28)-trienol and ergosta-5,7-dienol	0.17	1.75 1.19

* Not Detectable

TABLE 7

ERG3-ERG6 Mutants

<u>Sterol</u>	<u>Percent of Biomass</u>	
	<u>Non-transformed</u>	<u>Transformed</u>
	ATC6119	ATC2100 ATC2104
a. Squalene	N.D***	0.13 0.98
b. Zymosterol	0.21	1.10 1.80
c. Cholesta-7, 24-dienol	0.53	1.10 1.50

ERG5-ERG6 Mutants

<u>Sterol</u>	<u>Percent of Biomass</u>			
	<u>Non-transformed</u> ATC0402mu (n=4)	<u>Transformed</u>		
		ATC1503 (n=2)	ATC2401mu (n=4)	ATC1502 (n=2) ATC1500cp (n=1) ATC1551 (n=1)
a. Squalene	0.026	0.336	0.947	1.078 0.27 2.992
b. Zymosterol	1.107	1.358	1.125	2.065 3.746 5.125
c. C5,7,24*	1.542	0.956	1.064	1.354 1.868 2.372
d. C7,24**	0.213	0.362	0.250	0.408 0.564 0.775

* C5,7,24 is cholesta-5,7,24-trienol

** C7,24 is cholesta-7,24-dienol

*** Not Detectable

n = number of observations

The above data illustrate that transformation of mutants having a single defect in the expression of sterol biosynthetic pathway enzymes resulted in an increased accumulation of squalene and specific sterols (See Table 6).

Relative to a non-transformed erg3 mutant, erg3 mutants transformed with a plasmid vector useful in the present invention overaccumulated squalene, ergosta-8,22-dienol, ergosta-7,22-dienol, ergosta-8-enol and ergosta-7-enol.

Relative to a non-transformed erg5 mutant, erg5 mutants transformed with a plasmid vector useful in the present invention overaccumulated squalene, zymosterol, and a mixture of ergosta-5,7,24(28)-trienol and ergosta-5,7-dienol.

Similarly, transformation of mutants having double defects in the sterol biosynthetic pathway enzymes led to the overaccumulation of squalene and specific sterols.

Relative to a non-transformed erg3-erg6 mutant, erg3-erg6 mutants transformed with a plasmid vector useful in the present invention overaccumulated squalene, zymosterol and cholesta-7,24-dienol.

Relative to a non-transformed erg5-erg6 mutant, erg5-erg6 double mutants transformed with a plasmid vector useful in the present invention overaccumulated squalene, zymosterol, cholesta-5,7,24-trienol and cholesta-7,24-dienol.

The greatest increases in squalene and specific sterol accumulation are seen when erg5-erg6 mutant ATC0315rc is transformed with plasmid vector pARC304S (mutant ATC1551), as described in Example 9. Further, the data show that species ATC0402mu, the grandparent strain of mutant ATC0315rc, has elevated levels of sterols relative to either an erg5 or an erg6 single mutant (see Table 6).

EXAMPLE: 16 Induction of Squalene Accumulation in Yeast Transformant ATC1551

It is generally known that restricted aeration induces squalene accumulation at the expense of sterols in yeast cultures. This occurs because oxygen is required for the enzymatic conversion of squalene to squalene monoepoxide, which in turn is converted into lanosterol and other yeast sterols.

To determine if high levels of squalene accumulation could be induced in transformants, cultures of ATC1551 were grown under varying degrees of aeration by varying the volume (and hence the surface-to-volume ratio) of growth medium in shake-flask cultures and assaying squalene and total sterol at one day intervals over a period of four days.

Triplicate 250 ml baffled shake-flasks respectively containing 50, 100, 150 and 200 ml of YEP/2 percent glucose growth medium were inoculated with two ml of a 24 hour liquid culture of ATC1551 grown on a rotary shaker (200 rpm) at 30°C. Fifty ml culture aliquots were harvested by centrifugation after one, two, three and four days growth under the aforementioned conditions and lyophilized overnight.

To insure efficient squalene extraction, approximately 100 mg of each lyophilized sample was agitated for 10 minutes in 15 ml conical tubes containing a suitable quantity of glass beads and a small amount of water. The disrupted cell material was then extracted three successive times with 10 ml of 100 percent ethanol with vigorous agitation for one hour at 80°C. The combined ethanol extracts were reduced to dryness under a stream of nitrogen and redissolved in two ml of heptane containing 5 α -cholestane as the internal standard. GC analyses of squalene were conducted as previously described.

For total sterol analyses, the same samples were reduced under a stream of nitrogen and saponified in 5 ml of 95 percent ethanol/water solution containing 0.3 M KOH for one hour at 80°C. An equivalent volume of water was added and the samples were twice extracted with 10 ml aliquots of heptane. The heptane extracts were combined, reduced to a suitable volume and analyzed by GC.

The results are shown in Table 8 (data averaged from triplicate cultures and expressed as percent of dry biomass).

Table 8

		<u>Growth Medium Volume</u>			
		50 ml	100 ml	150 ml	200 ml
<u>Time to Harvest</u>		<u>Percent of Dry Biomass</u>			
5	Day 1				
	squalene	4.25	5.40	3.61	2.63
	total sterol	9.40	9.52	6.81	5.46
10	Day 2				
	squalene	4.78	6.43	11.89	8.32
	total sterol	8.29	6.44	3.72	2.98
15	Day 3				
	squalene	4.75	8.82	13.54	13.38
	total sterol	7.96	7.65	4.36	4.19
20	Day 4				
	squalene	4.03	7.08	15.99	14.72
	total sterol	7.09	8.62	5.10	3.39

The data show that in transformed, erg5-erg6 mutants, squalene is preferentially accumulated as compared to total sterol by restricting the level of aeration as compared to usual culture conditions (50 mls of growth medium), particularly after more than about one day of culture. The data also show that restricting the level of aeration (lowering the surface-to-volume ratio) also increases the sum total of squalene and total sterol accumulation, after more than about two days of culture.

Although the present invention has now been described in terms of certain preferred embodiments, and exemplified with respect thereto, one skilled in the art will readily appreciate that various modifications, changes, omissions and substitutions may be made without departing from the spirit thereof.

SEQUENCE LISTING

5

(1) GENERAL INFORMATION:

- 10 (i) APPLICANT: Saunders, Court A.
Wolf, Fred R.
Mukharji, Indrani
- (ii) TITLE OF INVENTION: A Method and Composition for Increasing
the Accumulation of Squalene and Specific Sterols in
15 Yeast
- (iii) NUMBER OF SEQUENCES: 24
- (iv) CORRESPONDENCE ADDRESS:
20 (A) ADDRESSEE: Amoco Corp., Patents and Licensing Dept.
(B) STREET: 200 East Randolph St.
(C) CITY: Chicago
(D) STATE: Illinois
(E) COUNTRY: USA
(F) ZIP: 60680-0703
- 25 (v) COMPUTER READABLE FORM:
(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.25
- 30 (vi) CURRENT APPLICATION DATA:
(A) APPLICATION NUMBER: 07/613,380
(B) FILING DATE: November 15, 1990
(C) CLASSIFICATION:
- 35 (viii) ATTORNEY/AGENT INFORMATION:
(A) NAME: Galloway, Norvall B.
- (ix) TELECOMMUNICATION INFORMATION:
(A) TELEPHONE: 312 856-7180
(B) TELEFAX: 312 856-4972

40

(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
45 (A) LENGTH: 3360 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

50

55

5

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 121..3282

10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

	TTTATTAACT TATTTTTTTC TTCTTTCTAC CCAATTCTAG TCAGGAAAAG ACTAAGGGCT	60
	GGAACATAGT GTATCATTGT CTAATTGTTG ATACAAAGTA GATAAATACA TAAAACAAGC	120
15	ATG CCG CCG CTA TTC AAG GGA CTG AAA CAG ATG GCA AAG CCA ATT GCC Met Pro Pro Leu Phe Lys Gly Leu Lys Gln Met Ala Lys Pro Ile Ala 1 5 10 15	168
	TAT GTT TCA AGA TTT TCG GCG AAA CGA CCA ATT CAT ATA ATA CTT TTT Tyr Val Ser Arg Phe Ser Ala Lys Arg Pro Ile His Ile Ile Leu Phe 20 25 30	216
20	TCT CTA ATC ATA TCC GCA TTC GCT TAT CTA TCC GTC ATT CAG TAT TAC Ser Leu Ile Ile Ser Ala Phe Ala Tyr Leu Ser Val Ile Gln Tyr Tyr 35 40 45	264
25	TTC AAT GGT TGG CAA CTA GAT TCA AAT AGT GTT TTT GAA ACT GCT CCA Phe Asn Gly Trp Gln Leu Asp Ser Asn Ser Val Phe Glu Thr Ala Pro 50 55 60	312
	AAT AAA GAC TCC AAC ACT CTA TTT CAA GAA TGT TCC CAT TAC TAC AGA Asn Lys Asp Ser Asn Thr Leu Phe Gln Glu Cys Ser His Tyr Tyr Arg 65 70 75 80	360
30	GAT TCC TCT CTA GAT GGT TGG GTA TCA ATC ACC GCG CAT GAA GCT AGT Asp Ser Ser Leu Asp Gly Trp Val Ser Ile Thr Ala His Glu Ala Ser 85 90 95	408
35	GAG TTA CCA GCC CCA CAC CAT TAC TAT CTA TTA AAC CTG AAC TTC AAT Glu Leu Pro Ala Pro His His Tyr Tyr Leu Leu Asn Leu Asn Phe Asn 100 105 110	456
	AGT CCT AAT GAA ACT GAC TCC ATT CCA GAA CTA GCT AAC ACG GTT TTT Ser Pro Asn Glu Thr Asp Ser Ile Pro Glu Leu Ala Asn Thr Val Phe 115 120 125	504
40	GAG AAA GAT AAT ACA AAA TAT ATT CTG CAA GAA GAT CTC AGT GTT TCC Glu Lys Asp Asn Thr Lys Tyr Ile Leu Gln Glu Asp Leu Ser Val Ser 130 135 140	552
45	AAA GAA ATT TCT TCT ACT GAT GGA ACG AAA TGG AGG TTA AGA AGT GAC Lys Glu Ile Ser Ser Thr Asp Gly Thr Lys Trp Arg Leu Arg Ser Asp 145 150 155 160	600

50

55

5

AGA AAA AGT CTT TTC GAC GTA AAG ACG TTA GCA TAT TCT CTC TAC GAT 648
 Arg Lys Ser Leu Phe Asp Val Lys Thr Leu Ala Tyr Ser Leu Tyr Asp
 165 170 175

10 GTA TTT TCA GAA AAT GTA ACC CAA GCA GAC CCG TTT GAC GTC CTT ATT 696
 Val Phe Ser Glu Asn Val Thr Gln Ala Asp Pro Phe Asp Val Leu Ile
 180 185 190

15 ATG GTT ACT GCC TAC CTA ATG ATG TTC TAC ACC ATA TTC GGC CTC TTC 744
 Met Val Thr Ala Tyr Leu Met Met Phe Tyr Thr Ile Phe Gly Leu Phe
 195 200 205

AAT GAC ATG AGG AAG ACC GGG TCA AAT TTT TGG TTG AGC GCC TCT ACA 792
 Asn Asp Met Arg Lys Thr Gly Ser Asn Phe Trp Leu Ser Ala Ser Thr
 210 215 220

20 GTG GTC AAT TCT GCA TCA TCA CTT TTC TTA GCA TTG TAT GTC ACC CAA 840
 Val Val Asn Ser Ala Ser Ser Leu Phe Leu Ala Leu Tyr Val Thr Gln
 225 230 235 240

TGT ATT CTA GGC AAA GAA GTT TCC GCA TTA ACT CTT TTT GAA GGT TTG 888
 Cys Ile Leu Gly Lys Glu Val Ser Ala Leu Thr Leu Phe Glu Gly Leu
 245 250 255

25 CCT TTC ATT GTA GTT GTT GTT GGT TTC AAG CAC AAA ATC AAG ATT GCC 936
 Pro Phe Ile Val Val Val Val Gly Phe Lys His Lys Ile Lys Ile Ala
 260 265 270

30 CAG TAT GCC CTG GAG AAA TTT GAA AGA GTC GGT TTA TCT AAA AGG ATT 984
 Gln Tyr Ala Leu Glu Lys Phe Glu Arg Val Gly Leu Ser Lys Arg Ile
 275 280 285

ACT ACC GAT GAA ATC GTT TTT GAA TCC GTG AGC GAA GAG GGT GGT CGT 1032
 Thr Thr Asp Glu Ile Val Phe Glu Ser Val Ser Glu Glu Gly Gly Arg
 290 295 300

35 TTG ATT CAA GAC CAT TTG CTT TGT ATT TTT GCC TTT ATC GGA TGC TCT 1080
 Leu Ile Gln Asp His Leu Leu Cys Ile Phe Ala Phe Ile Gly Cys Ser
 305 310 315 320

ATG TAT GCT CAC CAA TTG AAG ACT TTG ACA AAC TTC TGC ATA TTA TCA 1128
 Met Tyr Ala His Gln Leu Lys Thr Leu Thr Asn Phe Cys Ile Leu Ser
 325 330 335

40 GCA TTT ATC CTA ATT TTT GAA TTG ATT TTA ACT CCT ACA TTT TAT TCT 1176
 Ala Phe Ile Leu Ile Phe Glu Leu Ile Leu Thr Pro Thr Phe Tyr Ser
 340 345 350

45 GCT ATC TTA GCG CTT AGA CTG GAA ATG AAT GTT ATC CAC AGA TCT ACT 1224
 Ala Ile Leu Ala Leu Arg Leu Glu Met Asn Val Ile His Arg Ser Thr
 355 360 365

50

55

5

ATT ATC AAG CAA ACA TTA GAA GAA GAC GGT GTT GTT CCA TCT ACA GCA 1272
 Ile Ile Lys Gln Thr Leu Glu Glu Asp Gly Val Val Pro Ser Thr Ala
 370 375 380

10

AGA ATC ATT TCT AAA GCA GAA AAG AAA TCC GTA TCT TCT TTC TTA AAT 1320
 Arg Ile Ile Ser Lys Ala Glu Lys Lys Ser Val Ser Ser Ser Phe Leu Asn
 385 390 395 400

15

CTC AGT GTG GTT GTC ATT ATC ATG AAA CTC TCT GTC ATA CTG TTG TTT 1368
 Leu Ser Val Val Val Ile Ile Met Lys Leu Ser Val Ile Leu Leu Phe
 405 410 415

GTT TTC ATC AAC TTT TAT AAC TTT GGT GCA AAT TGG GTC AAT GAT GCC 1416
 Val Phe Ile Asn Phe Tyr Asn Phe Gly Ala Asn Trp Val Asn Asp Ala
 420 425 430

20

TTC AAT TCA TTG TAC TTC GAT AAG GAA CGT GTT TCT CTA CCA GAT TTT 1464
 Phe Asn Ser Leu Tyr Phe Asp Lys Glu Arg Val Ser Leu Pro Asp Phe
 435 440 445

ATT ACC TCG AAT GCC TCT GAA AAC TTT AAA GAG CAA GCT ATT GTT AGT 1512
 Ile Thr Ser Asn Ala Ser Glu Asn Phe Lys Glu Gln Ala Ile Val Ser
 450 455 460

25

GTC ACC CCA TTA TTA TAT TAC AAA CCC ATT AAG TCC TAC CAA CGC ATT 1560
 Val Thr Pro Leu Leu Tyr Tyr Lys Pro Ile Lys Ser Tyr Gln Arg Ile
 465 470 475 480

30

GAG GAT ATG GTT CTT CTA TTG CTT CGT AAT GTC AGT GTT GCC ATT CGT 1608
 Glu Asp Met Val Leu Leu Leu Leu Arg Asn Val Ser Val Ala Ile Arg
 485 490 495

GAT AGG TTC GTC AGT AAA TTA GTT CTT TCC GCC TTA GTA TGC AGT GCT 1656
 Asp Arg Phe Val Ser Lys Leu Val Leu Ser Ala Leu Val Cys Ser Ala
 500 505 510

35

GTC ATC AAT GTG TAT TTA TTG AAT GCT GCT AGA ATT CAT ACC AGT TAT 1704
 Val Ile Asn Val Tyr Leu Leu Asn Ala Ala Arg Ile His Thr Ser Tyr
 515 520 525

ACT GCA GAC CAA TTG GTG AAA ACT GAA GTC ACC AAG AAG TCT TTT ACT 1752
 Thr Ala Asp Gln Leu Val Lys Thr Glu Val Thr Lys Lys Ser Phe Thr
 530 535 540

40

GCT CCT GTA CAA AAG GCT TCT ACA CCA GTT TTA ACC AAT AAA ACA GTC 1800
 Ala Pro Val Gln Lys Ala Ser Thr Pro Val Leu Thr Asn Lys Thr Val
 545 550 555 560

45

ATT TCT GGA TCG AAA GTC AAA AGT TTA TCA TCT GCG CAA TCG AGC TCA 1848
 Ile Ser Gly Ser Lys Val Lys Ser Leu Ser Ser Ala Gln Ser Ser Ser
 565 570 575

50

55

5

TCA GGA CCT TCA TCA TCT AGT GAG GAA GAT GAT TCC CGC GAT ATT GAA 1896
 Ser Gly Pro Ser Ser Ser Ser Glu Glu Asp Asp Ser Arg Asp Ile Glu
 580 585 590

10 AGC TTG GAT AAG AAA ATA CGT CCT TTA GAA GAA TTA GAA GCA TTA TTA 1944
 Ser Leu Asp Lys Lys Ile Arg Pro Leu Glu Glu Leu Glu Ala Leu Leu
 595 600 605

15 AGT AGT GGA AAT ACA AAA CAA TTG AAG AAC AAA GAG GTC GCT GCC TTG 1992
 Ser Ser Gly Asn Thr Lys Lys Leu Lys Asn Lys Glu Val Ala Ala Leu
 610 615 620

GTT ATT CAC GGT AAG TTA CCT TTG TAC GCT TTG GAG AAA AAA TTA GGT 2040
 Val Ile His Gly Lys Leu Pro Leu Tyr Ala Leu Lys Lys Leu Gly
 625 630 635 640

20 GAT ACT ACG AGA GCG GTT GCG GTA CGT AGG AAG GCT CTT TCA ATT TTG 2088
 Asp Thr Thr Arg Ala Val Ala Val Arg Arg Lys Ala Leu Ser Ile Leu
 645 650 655

GCA GAA GCT CCT GTA TTA GCA TCT GAT CGT TTA CCA TAT AAA AAT TAT 2136
 Ala Glu Ala Pro Val Leu Ala Ser Asp Arg Leu Pro Tyr Lys Asn Tyr
 660 665 670

25 GAC TAC GAC CGC GTA TTT GGC GCT TGT TGT GAA AAT GTT ATA GGT TAC 2184
 Asp Tyr Asp Arg Val Phe Gly Ala Cys Cys Glu Asn Val Ile Gly Tyr
 675 680 685

30 ATG CCT TTG CCC GTT GGT GTT ATA GGC CCC TTG GTT ATC GAT GGT ACA 2232
 Met Pro Leu Pro Val Gly Val Ile Gly Pro Leu Val Ile Asp Gly Thr
 690 695 700

TCT TAT CAT ATA CCA ATG GCA ACT ACA GAG GGT TGT TTG GTA GCT TCT 2280
 Ser Tyr His Ile Pro Met Ala Thr Thr Glu Gly Cys Leu Val Ala Ser
 705 710 715 720

35 GCC ATG CGT GGC TGT AAG GCA ATC AAT GCT GGC GGT GGT GCA ACA ACT 2328
 Ala Met Arg Gly Cys Lys Ala Ile Asn Ala Gly Gly Gly Ala Thr Thr
 725 730 735

GTT TTA ACT AAG GAT GGT ATG ACA AGA GGC CCA GTA GTC CGT TTC CCA 2376
 Val Leu Thr Lys Asp Gly Met Thr Arg Gly Pro Val Val Arg Phe Pro
 740 745 750

40 ACT TTG AAA AGA TCT GGT GCC TGT AAG ATA TGG TTA GAC TCA GAA GAG 2424
 Thr Leu Lys Arg Ser Gly Ala Cys Lys Ile Trp Leu Asp Ser Glu Glu
 755 760 765

45 GGA CAA AAC GCA ATT AAA AAA GCT TTT AAC TCT ACA TCA AGA TTT GCA 2472
 Gly Gln Asn Ala Ile Lys Lys Ala Phe Asn Ser Thr Ser Arg Phe Ala
 770 775 780

50

55

5	CGT	CTG	CAA	CAT	ATT	CAA	ACT	TGT	CTA	GCA	GGA	GAT	TTA	CTC	TTC	ATG	2520
	Arg	Leu	Gln	His	Ile	Gln	Thr	Cys	Leu	Ala	Gly	Asp	Leu	Leu	Phe	Met	
	785					790					795					800	
10	AGA	TTT	AGA	ACA	ACT	ACT	GGT	GAC	GCA	ATG	GGT	ATG	AAT	ATG	ATT	TCT	2568
	Arg	Phe	Arg	Thr	Thr	Thr	Gly	Asp	Ala	Met	Gly	Met	Asn	Met	Ile	Ser	
					805					810					815		
	AAA	GGT	GTC	GAA	TAC	TCA	TTA	AAG	CAA	ATG	GTA	GAA	GAG	TAT	GGC	TGG	2616
	Lys	Gly	Val	Glu	Tyr	Ser	Leu	Lys	Gln	Met	Val	Glu	Glu	Tyr	Gly	Trp	
				820					825					830			
15	GAA	GAT	ATG	GAG	GTT	GTC	TCC	GTT	TCT	GGT	AAC	TAC	TGT	ACC	GAC	AAA	2664
	Glu	Asp	Met	Glu	Val	Val	Ser	Val	Ser	Gly	Asn	Tyr	Cys	Thr	Asp	Lys	
			835					840					845				
20	AAA	CCA	GCT	GCC	ATC	AAC	TGG	ATC	GAA	GGT	CGT	GGT	AAG	AGT	GTC	GTC	2712
	Lys	Pro	Ala	Ala	Ile	Asn	Trp	Ile	Glu	Gly	Arg	Gly	Lys	Ser	Val	Val	
		850					855					860					
	GCA	GAA	GCT	ACT	ATT	CCT	GGT	GAT	GTT	GTC	AGA	AAA	GTG	TTA	AAA	AGT	2760
	Ala	Glu	Ala	Thr	Ile	Pro	Gly	Asp	Val	Val	Arg	Lys	Val	Leu	Lys	Ser	
	865					870					875					880	
25	GAT	GTT	TCC	GCA	TTG	GTT	GAG	TTG	AAC	ATT	GCT	AAG	AAT	TTG	GTT	GGA	2808
	Asp	Val	Ser	Ala	Leu	Val	Glu	Leu	Asn	Ile	Ala	Lys	Asn	Leu	Val	Gly	
					885					890					895		
	TCT	GCA	ATG	GCT	GGG	TCT	GTT	GGT	GGA	TTT	AAC	GCA	CAT	GCA	GCT	AAT	2856
	Ser	Ala	Met	Ala	Gly	Ser	Val	Gly	Gly	Phe	Asn	Ala	His	Ala	Ala	Asn	
30				900					905					910			
	TTA	GTG	ACA	GCT	GTT	TTC	TTG	GCA	TTA	GGA	CAA	GAT	CCT	GCA	CAA	AAT	2904
	Leu	Val	Thr	Ala	Val	Phe	Leu	Ala	Leu	Gly	Gln	Asp	Pro	Ala	Gln	Asn	
			915					920					925				
35	GTT	GAA	AGT	TCC	AAC	TGT	ATA	ACA	TTG	ATG	AAA	GAA	GTG	GAC	GGT	GAT	2952
	Val	Glu	Ser	Ser	Asn	Cys	Ile	Thr	Leu	Met	Lys	Glu	Val	Asp	Gly	Asp	
		930				935						940					
	TTG	AGA	ATT	TCC	GTA	TCC	ATG	CCA	TCC	ATC	GAA	GTA	GGT	ACC	ATC	GGT	3000
	Leu	Arg	Ile	Ser	Val	Ser	Met	Pro	Ser	Ile	Glu	Val	Gly	Thr	Ile	Gly	
	945					950					955					960	
40	GGT	GGT	ACT	GTT	CTA	GAA	CCA	CAA	GGT	GCC	ATG	TTG	GAC	TTA	TTA	GGT	3048
	Gly	Gly	Thr	Val	Leu	Glu	Pro	Gln	Gly	Ala	Met	Leu	Asp	Leu	Leu	Gly	
					965				970						975		
45	GTA	AGA	GGC	CCG	CAT	GCT	ACC	GCT	CCT	GGT	ACC	AAC	GCA	CGT	CAA	TTA	3096
	Val	Arg	Gly	Pro	His	Ala	Thr	Ala	Pro	Gly	Thr	Asn	Ala	Arg	Gln	Leu	
				980					985					990			

50

55

5

GCA AGA ATA GTT GCC TGT GCC GTC TTG GCA GGT GAA TTA TCC TTA TGT 3144
 Ala Arg Ile Val Ala Cys Ala Val Leu Ala Gly Glu Leu Ser Leu Cys
 995 1000 1005
 10 GCT GCC CTA GCA GCC GGC CAT TTG GTT CAA AGT CAT ATG ACC CAC AAC 3192
 Ala Ala Leu Ala Ala Gly His Leu Val Gln Ser His Met Thr His Asn
 1010 1015 1020
 AGG AAA CCT GCT GAA CCA ACA AAA CCT AAC AAT TTG GAC GCC ACT GAT 3240
 Arg Lys Pro Ala Glu Pro Thr Lys Pro Asn Asn Leu Asp Ala Thr Asp
 1025 1030 1035 1040
 15 ATA AAT CGT TTG AAA GAT GGG TCC GTC ACC TGC ATT AAA TCC 3282
 Ile Asn Arg Leu Lys Asp Gly Ser Val Thr Cys Ile Lys Ser
 1045 1050
 TAAACTTAGT CATACGTCAT TGGTATTCTC TTGAAAAAGA AGCACAACAG CACCATGTGT 3342
 20 TACGTAAAAT ATTACTT 3360

(2) INFORMATION FOR SEQ ID NO:2:

25

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 1054 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

30

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met Pro Pro Leu Phe Lys Gly Leu Lys Gln Met Ala Lys Pro Ile Ala
 1 5 10 15
 Tyr Val Ser Arg Phe Ser Ala Lys Arg Pro Ile His Ile Ile Leu Phe
 20 25 30
 35 Ser Leu Ile Ile Ser Ala Phe Ala Tyr Leu Ser Val Ile Gln Tyr Tyr
 35 40 45
 Phe Asn Gly Trp Gln Leu Asp Ser Asn Ser Val Phe Glu Thr Ala Pro
 50 55 60
 40 Asn Lys Asp Ser Asn Thr Leu Phe Gln Glu Cys Ser His Tyr Tyr Arg
 65 70 75 80
 Asp Ser Ser Leu Asp Gly Trp Val Ser Ile Thr Ala His Glu Ala Ser
 85 90 95
 45 Glu Leu Pro Ala Pro His His Tyr Tyr Leu Leu Asn Leu Asn Phe Asn
 100 105 110

50

55

5 Ser Pro Asn Glu Thr Asp Ser Ile Pro Glu Leu Ala Asn Thr Val Phe
 115 120 125
 Glu Lys Asp Asn Thr Lys Tyr Ile Leu Gln Glu Asp Leu Ser Val Ser
 130 135 140
 10 Lys Glu Ile Ser Ser Thr Asp Gly Thr Lys Trp Arg Leu Arg Ser Asp
 145 150 155 160
 Arg Lys Ser Leu Phe Asp Val Lys Thr Leu Ala Tyr Ser Leu Tyr Asp
 165 170 175
 15 Val Phe Ser Glu Asn Val Thr Gln Ala Asp Pro Phe Asp Val Leu Ile
 180 185 190
 Met Val Thr Ala Tyr Leu Met Met Phe Tyr Thr Ile Phe Gly Leu Phe
 195 200 205
 20 Asn Asp Met Arg Lys Thr Gly Ser Asn Phe Trp Leu Ser Ala Ser Thr
 210 215 220
 Val Val Asn Ser Ala Ser Ser Leu Phe Leu Ala Leu Tyr Val Thr Gln
 225 230 235 240
 25 Cys Ile Leu Gly Lys Glu Val Ser Ala Leu Thr Leu Phe Glu Gly Leu
 245 250 255
 Pro Phe Ile Val Val Val Val Gly Phe Lys His Lys Ile Lys Ile Ala
 260 265 270
 30 Gln Tyr Ala Leu Glu Lys Phe Glu Arg Val Gly Leu Ser Lys Arg Ile
 275 280 285
 Thr Thr Asp Glu Ile Val Phe Glu Ser Val Ser Glu Glu Gly Gly Arg
 290 295 300
 35 Leu Ile Gln Asp His Leu Leu Cys Ile Phe Ala Phe Ile Gly Cys Ser
 305 310 315 320
 40 Met Tyr Ala His Gln Leu Lys Thr Leu Thr Asn Phe Cys Ile Leu Ser
 325 330 335
 Ala Phe Ile Leu Ile Phe Glu Leu Ile Leu Thr Pro Thr Phe Tyr Ser
 340 345 350
 45 Ala Ile Leu Ala Leu Arg Leu Glu Met Asn Val Ile His Arg Ser Thr
 355 360 365
 Ile Ile Lys Gln Thr Leu Glu Glu Asp Gly Val Val Pro Ser Thr Ala
 370 375 380

50

55

5 Arg Ile Ile Ser Lys Ala Glu Lys Lys Ser Val Ser Ser Phe Leu Asn
 385 390 395 400
 Leu Ser Val Val Val Ile Ile Met Lys Leu Ser Val Ile Leu Leu Phe
 405 410 415
 10 Val Phe Ile Asn Phe Tyr Asn Phe Gly Ala Asn Trp Val Asn Asp Ala
 420 425 430
 Phe Asn Ser Leu Tyr Phe Asp Lys Glu Arg Val Ser Leu Pro Asp Phe
 435 440 445
 15 Ile Thr Ser Asn Ala Ser Glu Asn Phe Lys Glu Gln Ala Ile Val Ser
 450 455 460
 Val Thr Pro Leu Leu Tyr Tyr Lys Pro Ile Lys Ser Tyr Gln Arg Ile
 465 470 475 480
 20 Glu Asp Met Val Leu Leu Leu Leu Arg Asn Val Ser Val Ala Ile Arg
 485 490 495
 Asp Arg Phe Val Ser Lys Leu Val Leu Ser Ala Leu Val Cys Ser Ala
 500 505 510
 25 Val Ile Asn Val Tyr Leu Leu Asn Ala Ala Arg Ile His Thr Ser Tyr
 515 520 525
 Thr Ala Asp Gln Leu Val Lys Thr Glu Val Thr Lys Lys Ser Phe Thr
 530 535 540
 30 Ala Pro Val Gln Lys Ala Ser Thr Pro Val Leu Thr Asn Lys Thr Val
 545 550 555 560
 Ile Ser Gly Ser Lys Val Lys Ser Leu Ser Ser Ala Gln Ser Ser Ser
 565 570 575
 35 Ser Gly Pro Ser Ser Ser Ser Glu Glu Asp Asp Ser Arg Asp Ile Glu
 580 585 590
 40 Ser Leu Asp Lys Lys Ile Arg Pro Leu Glu Glu Leu Glu Ala Leu Leu
 595 600 605
 Ser Ser Gly Asn Thr Lys Gln Leu Lys Asn Lys Glu Val Ala Ala Leu
 610 615 620
 45 Val Ile His Gly Lys Leu Pro Leu Tyr Ala Leu Glu Lys Lys Leu Gly
 625 630 635 640
 Asp Thr Thr Arg Ala Val Ala Val Arg Arg Lys Ala Leu Ser Ile Leu
 645 650 655
 50 Ala Glu Ala Pro Val Leu Ala Ser Asp Arg Leu Pro Tyr Lys Asn Tyr

55

	660					665					670					
5	Asp	Tyr	Asp	Arg	Val	Phe	Gly	Ala	Cys	Cys	Glu	Asn	Val	Ile	Gly	Tyr
			675					680					685			
10	Met	Pro	Leu	Pro	Val	Gly	Val	Ile	Gly	Pro	Leu	Val	Ile	Asp	Gly	Thr
		690					695					700				
15	Ser	Tyr	His	Ile	Pro	Met	Ala	Thr	Thr	Glu	Gly	Cys	Leu	Val	Ala	Ser
	705					710					715					720
	Ala	Met	Arg	Gly	Cys	Lys	Ala	Ile	Asn	Ala	Gly	Gly	Gly	Ala	Thr	Thr
					725					730					735	
	Val	Leu	Thr	Lys	Asp	Gly	Met	Thr	Arg	Gly	Pro	Val	Val	Arg	Phe	Pro
				740					745					750		
20	Thr	Leu	Lys	Arg	Ser	Gly	Ala	Cys	Lys	Ile	Trp	Leu	Asp	Ser	Glu	Glu
			755					760					765			
	Gly	Gln	Asn	Ala	Ile	Lys	Lys	Ala	Phe	Asn	Ser	Thr	Ser	Arg	Phe	Ala
		770					775					780				
25	Arg	Leu	Gln	His	Ile	Gln	Thr	Cys	Leu	Ala	Gly	Asp	Leu	Leu	Phe	Met
	785					790					795					800
	Arg	Phe	Arg	Thr	Thr	Thr	Gly	Asp	Ala	Met	Gly	Met	Asn	Met	Ile	Ser
					805					810					815	
30	Lys	Gly	Val	Glu	Tyr	Ser	Leu	Lys	Gln	Met	Val	Glu	Glu	Tyr	Gly	Trp
				820					825					830		
	Glu	Asp	Met	Glu	Val	Val	Ser	Val	Ser	Gly	Asn	Tyr	Cys	Thr	Asp	Lys
			835					840					845			
35	Lys	Pro	Ala	Ala	Ile	Asn	Trp	Ile	Glu	Gly	Arg	Gly	Lys	Ser	Val	Val
		850					855					860				
	Ala	Glu	Ala	Thr	Ile	Pro	Gly	Asp	Val	Val	Arg	Lys	Val	Leu	Lys	Ser
	865					870					875					880
40	Asp	Val	Ser	Ala	Leu	Val	Glu	Leu	Asn	Ile	Ala	Lys	Asn	Leu	Val	Gly
					885					890					895	
	Ser	Ala	Met	Ala	Gly	Ser	Val	Gly	Gly	Phe	Asn	Ala	His	Ala	Ala	Asn
				900					905					910		
45	Leu	Val	Thr	Ala	Val	Phe	Leu	Ala	Leu	Gly	Gln	Asp	Pro	Ala	Gln	Asn
			915					920					925			
	Val	Glu	Ser	Ser	Asn	Cys	Ile	Thr	Leu	Met	Lys	Glu	Val	Asp	Gly	Asp
		930					935					940				

55

5

Leu Arg Ile Ser Val Ser Met Pro Ser Ile Glu Val Gly Thr Ile Gly
945 950 955 960

10

Gly Gly Thr Val Leu Glu Pro Gln Gly Ala Met Leu Asp Leu Leu Gly
965 970 975

Val Arg Gly Pro His Ala Thr Ala Pro Gly Thr Asn Ala Arg Gln Leu
980 985 990

15

Ala Arg Ile Val Ala Cys Ala Val Leu Ala Gly Glu Leu Ser Leu Cys
995 1000 1005

Ala Ala Leu Ala Ala Gly His Leu Val Gln Ser His Met Thr His Asn
1010 1015 1020

Arg Lys Pro Ala Glu Pro Thr Lys Pro Asn Asn Leu Asp Ala Thr Asp
1025 1030 1035 1040

20

Ile Asn Arg Leu Lys Asp Gly Ser Val Thr Cys Ile Lys Ser
1045 1050

(2) INFORMATION FOR SEQ ID NO:3:

25

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 4768 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

30

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 164..2827

35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

TGTATGTCTT GTCTTTCTCC TAAGGGGCGT AGGCTCATTG ATAACATCATG TCCTCACCTT 60

GCACTCCTTT TGGAATTATT TGGTTTGAGT GAAGAAGACC GGACCTTCGA GGTTCGCAAC 120

40

TTAAACAATA GACTTGTGAG GATCCAGGGA CCGAGTGGCT ACA ATG TTG TCA CGA 175
Met Leu Ser Arg
1

45

CTT TTC CGT ATG CAT GGC CTC TTT GTG GCC TCC CAT CCC TGG GAA GTT 223
Leu Phe Arg Met His Gly Leu Phe Val Ala Ser His Pro Trp Glu Val
5 10 15 20

50

55

5

ATT GTG GGG ACG GTG ACA CTT ACC ATC TGT ATG ATG TCC ATG AAC ATG 271
 Ile Val Gly Thr Val Thr Leu Thr Ile Cys Met Met Ser Met Asn Met
 25 30 35

10

TTC ACT GGC AAC AAC AAG ATC TGT GGT TGG AAT TAC GAG TGC CCA AAA 319
 Phe Thr Gly Asn Asn Lys Ile Cys Gly Trp Asn Tyr Glu Cys Pro Lys
 40 45 50

15

TTT GAG GAG GAT GTA TTG AGC AGT GAC ATC ATC ATC CTC ACC ATA ACA 367
 Phe Glu Glu Asp Val Leu Ser Ser Asp Ile Ile Ile Leu Thr Ile Thr
 55 60 65

CGG TGC ATC GCC ATC CTG TAC ATT TAC TTC CAG TTC CAG AAC TTA CGT 415
 Arg Cys Ile Ala Ile Leu Tyr Ile Tyr Phe Gln Phe Gln Asn Leu Arg
 70 75 80

20

CAG CTT GGG TCG AAG TAT ATT TTA GGT ATT GCT GGC CTG TTC ACA ATT 463
 Gln Leu Gly Ser Lys Tyr Ile Leu Gly Ile Ala Gly Leu Phe Thr Ile
 85 90 95 100

25

TTC TCA AGT TTT GTC TTT AGT ACA GTC GTC ATT CAC TTC TTA GAC AAA 511
 Phe Ser Ser Phe Val Phe Ser Thr Val Val Ile His Phe Leu Asp Lys
 105 110 115

GAA CTG ACG GGC TTA AAT GAA GCT TTG CCC TTT TTC CTG CTT TTG ATT 559
 Glu Leu Thr Gly Leu Asn Glu Ala Leu Pro Phe Phe Leu Leu Leu Ile
 120 125 130

30

GAC CTT TCT AGA GCG AGT GCA CTA GCA AAG TTT GCC CTA AGT TCA AAC 607
 Asp Leu Ser Arg Ala Ser Ala Leu Ala Lys Phe Ala Leu Ser Ser Asn
 135 140 145

TCT CAG GAT GAA GTA AGG GAA AAT ATA GCT CGC GGA ATG GCA ATT CTG 655
 Ser Gln Asp Glu Val Arg Glu Asn Ile Ala Arg Gly Met Ala Ile Leu
 150 155 160

35

GGC CCC ACA TTC ACC CTT GAT GCT CTT GTG GAA TGT CTT GTA ATT GGA 703
 Gly Pro Thr Phe Thr Leu Asp Ala Leu Val Glu Cys Leu Val Ile Gly
 165 170 175 180

40

GTT GGC ACC ATG TCA GGG GTG CGT CAG CTT GAA ATC ATG TGC TGC TTT 751
 Val Gly Thr Met Ser Gly Val Arg Gln Leu Glu Ile Met Cys Cys Phe
 185 190 195

GGC TGC ATG TCT GTG CTT GCC AAC TAC TTC GTG TTC ATG ACA TTT TTC 799
 Gly Cys Met Ser Val Leu Ala Asn Tyr Phe Val Phe Met Thr Phe Phe
 200 205 210

45

CCA GCG TGT GTG TCC CTG GTC CTT GAG CTT TCT CGG GAA AGT CGA GAG 847
 Pro Ala Cys Val Ser Leu Val Leu Glu Leu Ser Arg Glu Ser Arg Glu
 215 220 225

50

55

5	GGT CGT CCA ATT TGG CAG CTT AGC CAT TTT GCC CGA GTT TTG GAA GAA Gly Arg Pro Ile Trp Gln Leu Ser His Phe Ala Arg Val Leu Glu Glu	895
	230 235 240	
10	GAA GAG AAT AAA CCA AAC CCT GTA ACC CAA AGG GTC AAG ATG ATT ATG Glu Glu Asn Lys Pro Asn Pro Val Thr Gln Arg Val Lys Met Ile Met	943
	245 250 255 260	
	TCT TTA GGT TTG GTT CTT GTT CAT GCT CAC AGT CGA TGG ATA GCT GAT Ser Leu Gly Leu Val Leu Val His Ala His Ser Arg Trp Ile Ala Asp	991
	265 270 275	
15	CCT TCC CCT CAG AAT AGC ACA ACA GAA CAT TCT AAA GTC TCC TTG GGA Pro Ser Pro Gln Asn Ser Thr Thr Glu His Ser Lys Val Ser Leu Gly	1039
	280 285 290	
20	CTG GAT GAA GAT GTG TCC AAG AGA ATT GAA CCA AGT GTT TCT CTC TGG Leu Asp Glu Asp Val Ser Lys Arg Ile Glu Pro Ser Val Ser Leu Trp	1087
	295 300 305	
	CAG TTT TAT CTC TCC AAG ATG ATC AGC ATG GAC ATT GAA CAA GTG GTT Gln Phe Tyr Leu Ser Lys Met Ile Ser Met Asp Ile Glu Gln Val Val	1135
	310 315 320	
25	ACC CTG AGC TTA GCT TTT CTG TTG GCT GTC AAG TAC ATT TTC TTT GAA Thr Leu Ser Leu Ala Phe Leu Leu Ala Val Lys Tyr Ile Phe Phe Glu	1183
	325 330 335 340	
	CAA GCA GAG ACA GAG TCC ACA CTG TCT TTA AAA AAT CCT ATC ACG TCT Gln Ala Glu Thr Glu Ser Thr Leu Ser Leu Lys Asn Pro Ile Thr Ser	1231
	345 350 355	
30	CCT GTC GTG ACC CCA AAG AAA GCT CCA GAC AAC TGT TGT AGA CGG GAG Pro Val Val Thr Pro Lys Lys Ala Pro Asp Asn Cys Cys Arg Arg Glu	1279
	360 365 370	
35	CCT CTG CTT GTG AGA AGG AGC GAG AAG CTT TCA TCG GTT GAG GAG GAG Pro Leu Leu Val Arg Arg Ser Glu Lys Leu Ser Ser Val Glu Glu Glu	1327
	375 380 385	
	CCT GGG GTG AGC CAA GAT AGA AAA GTT GAG GTT ATA AAA CCA TTA GTG Pro Gly Val Ser Gln Asp Arg Lys Val Glu Val Ile Lys Pro Leu Val	1375
	390 395 400	
40	GTG GAA ACT GAG AGT GCA AGC AGA GCT ACA TTT GTG CTT GGC GCC TCT Val Glu Thr Glu Ser Ala Ser Arg Ala Thr Phe Val Leu Gly Ala Ser	1423
	405 410 415 420	
45	GGG ACC AGC CCT CCA GTG GCA GCG AGG ACA CAG GAG CTT GAA ATT GAA Gly Thr Ser Pro Pro Val Ala Ala Arg Thr Gln Glu Leu Glu Ile Glu	1471
	425 430 435	

50

55

5

10

15

20

25

30

35

40

45

50

55

CTC	CCC	AGT	GAG	CCT	CGG	CCT	AAT	GAA	GAA	TGT	CTG	CAG	ATA	CTG	GAG	1519
Leu	Pro	Ser	Glu	Pro	Arg	Pro	Asn	Glu	Glu	Cys	Leu	Gln	Ile	Leu	Glu	
			440					445					450			
AGT	GCC	GAG	AAA	GGT	GCA	AAG	TTC	CTT	AGC	GAT	GCA	GAG	ATC	ATC	CAG	1567
Ser	Ala	Glu	Lys	Gly	Ala	Lys	Phe	Leu	Ser	Asp	Ala	Glu	Ile	Ile	Gln	
			455				460					465				
TTG	GTC	AAT	GCC	AAG	CAC	ATC	CCA	GCC	TAC	AAA	TTG	GAA	ACC	TTA	ATG	1615
Leu	Val	Asn	Ala	Lys	His	Ile	Pro	Ala	Tyr	Lys	Leu	Glu	Thr	Leu	Met	
			470			475					480					
GAA	ACT	CAT	GAA	CGT	GGT	GTA	TCT	ATT	CGC	CGG	CAG	CTC	CTC	TCC	ACA	1663
Glu	Thr	His	Glu	Arg	Gly	Val	Ser	Ile	Arg	Arg	Gln	Leu	Leu	Ser	Thr	
			485		490				495						500	
AAG	CTT	CCA	GAG	CCT	TCT	TCT	CTG	CAG	TAC	CTG	CCT	TAC	AGA	GAT	TAT	1711
Lys	Leu	Pro	Glu	Pro	Ser	Ser	Leu	Gln	Tyr	Leu	Pro	Tyr	Arg	Asp	Tyr	
				505					510					515		
AAT	TAT	TCC	CTG	GTG	ATG	GGA	GCT	TGC	TGT	GAG	AAT	GTG	ATC	GGA	TAT	1759
Asn	Tyr	Ser	Leu	Val	Met	Gly	Ala	Cys	Cys	Glu	Asn	Val	Ile	Gly	Tyr	
			520					525					530			
ATG	CCC	ATC	CCT	GTC	GGA	GTA	GCA	GGG	CCT	CTG	TGC	CTG	GAT	GGT	AAA	1807
Met	Pro	Ile	Pro	Val	Gly	Val	Ala	Gly	Pro	Leu	Cys	Leu	Asp	Gly	Lys	
			535				540					545				
GAG	TAC	CAG	GTT	CCA	ATG	GCA	ACA	ACG	GAA	GGC	TGT	CTG	GTG	GCC	AGC	1855
Glu	Tyr	Gln	Val	Pro	Met	Ala	Thr	Thr	Glu	Gly	Cys	Leu	Val	Ala	Ser	
			550			555					560					
ACC	AAC	AGA	GGC	TGC	AGG	GCA	ATA	GGT	CTT	GGT	GGA	GGT	GCC	AGC	AGC	1903
Thr	Asn	Arg	Gly	Cys	Arg	Ala	Ile	Gly	Leu	Gly	Gly	Gly	Ala	Ser	Ser	
			565		570				575						580	
CGG	GTC	CTT	GCA	GAT	GGG	ATG	ACC	CGG	GGC	CCA	GTG	GTG	CGT	CTT	CCT	1951
Arg	Val	Leu	Ala	Asp	Gly	Met	Thr	Arg	Gly	Pro	Val	Val	Arg	Leu	Pro	
				585				590						595		
CGT	GCT	TGT	GAT	TCT	GCA	GAA	GTG	AAG	GCC	TGG	CTT	GAA	ACA	CCC	GAA	1999
Arg	Ala	Cys	Asp	Ser	Ala	Glu	Val	Lys	Ala	Trp	Leu	Glu	Thr	Pro	Glu	
			600					605					610			
GGG	TTT	GCG	GTG	ATA	AAG	GAC	GCC	TTC	GAT	AGC	ACT	AGC	AGA	TTT	GCA	2047
Gly	Phe	Ala	Val	Ile	Lys	Asp	Ala	Phe	Asp	Ser	Thr	Ser	Arg	Phe	Ala	
			615				620					625				
CGT	CTA	CAG	AAG	CTT	CAT	GTG	ACC	ATG	GCA	GGG	CGC	AAC	CTG	TAC	ATC	2095
Arg	Leu	Gln	Lys	Leu	His	Val	Thr	Met	Ala	Gly	Arg	Asn	Leu	Tyr	Ile	
			630			635					640					

5		CGT TTC CAG TCC AAG ACA GGG GAT GCC ATG GGG ATG AAC ATG ATT TCC	2143
		Arg Phe Gln Ser Lys Thr Gly Asp Ala Met Gly Met Asn Met Ile Ser	
		645 650 655 660	
10		AAG GGC ACT GAG AAA GCA CTT CTG AAG CTT CAG GAG TTC TTT CCT GAA	2191
		Lys Gly Thr Glu Lys Ala Leu Leu Lys Leu Gln Glu Phe Phe Pro Glu	
		665 670 675	
		ATG CAG ATT CTG GCA GTT AGT GGT AAC TAC TGC ACT GAC AAG AAA CCT	2239
		Met Gln Ile Leu Ala Val Ser Gly Asn Tyr Cys Thr Asp Lys Lys Pro	
		680 685 690	
15		GCC GCC ATA AAC TGG ATC GAG GGA AGA GGA AAG ACA GTT GTG TGT GAA	2287
		Ala Ala Ile Asn Trp Ile Glu Gly Arg Gly Lys Thr Val Val Cys Glu	
		695 700 705	
20		GCT GTT ATT CCA GCC AAG GTG GTG AGA GAA GTA TTA AAG ACA ACT ACG	2335
		Ala Val Ile Pro Ala Lys Val Val Arg Glu Val Leu Lys Thr Thr Thr	
		710 715 720	
		GAA GCT ATG ATT GAC GTA AAC ATT AAC AAG AAT CTT GTG GGT TCT GCC	2383
		Glu Ala Met Ile Asp Val Asn Ile Asn Lys Asn Leu Val Gly Ser Ala	
		725 730 735 740	
25		ATG GCT GGG AGC ATA GGA GGC TAC AAT GCC CAT GCA GCA AAC ATC GTC	2431
		Met Ala Gly Ser Ile Gly Gly Tyr Asn Ala His Ala Ala Asn Ile Val	
		745 750 755	
		ACT GCT ATC TAC ATT GCA TGT GGC CAG GAT GCA GCA CAG AAT GTG GGG	2479
		Thr Ala Ile Tyr Ile Ala Cys Gly Gln Asp Ala Ala Gln Asn Val Gly	
		760 765 770	
30		AGT TCA AAC TGT ATT ACT TTA ATG GAA GCA AGT GGT CCC ACG AAT GAA	2527
		Ser Ser Asn Cys Ile Thr Leu Met Glu Ala Ser Gly Pro Thr Asn Glu	
		775 780 785	
35		GAC TTG TAT ATC AGC TGC ACC ATG CCA TCT ATA GAG ATA GGA ACT GTG	2575
		Asp Leu Tyr Ile Ser Cys Thr Met Pro Ser Ile Glu Ile Gly Thr Val	
		790 795 800	
		GGT GGT GGG ACC AAC CTC CTA CCA CAG CAG GCC TGT CTG CAG ATG CTA	2623
		Gly Gly Gly Thr Asn Leu Leu Pro Gln Gln Ala Cys Leu Gln Met Leu	
		805 810 815 820	
40		GGT GTT CAA GGA GCG TGC AAA GAC AAT CCT GGA GAA AAT GCA CGG CAA	2671
		Gly Val Gln Gly Ala Cys Lys Asp Asn Pro Gly Glu Asn Ala Arg Gln	
		825 830 835	
45		CTT GCC CGA ATT GTG TGT GGT ACT GTA ATG GCT GGG GAG TTG TCC TTG	2719
		Leu Ala Arg Ile Val Cys Gly Thr Val Met Ala Gly Glu Leu Ser Leu	
		840 845 850	

50

55

5
 ATG GCA GCA TTG GCA GCA GGA CAT CTT GTT AGA AGT CAC ATG GTT CAT 2767
 Met Ala Ala Leu Ala Ala Gly His Leu Val Arg Ser His Met Val His
 855 860 865

10
 AAC AGA TCG AAG ATA AAT TTA CAA GAT CTG CAA GGA ACG TGC ACC AAG 2815
 Asn Arg Ser Lys Ile Asn Leu Gln Asp Leu Gln Gly Thr Cys Thr Lys
 870 875 880

AAG TCA GCT TGAGCAGCCT GACAGTATTG AACTGAAACA CGGGCATTGG 2864
 Lys Ser Ala
 885

15
 GTTCTCAAGG ACTAACATGA AATCTGTGAA TTAAAAATCT CAATGCAGTG TCTTGTGGAA 2924
 GATGAATGAA CGTGATCAGT GAGACGCCTG CTTGGTTTCT GGCTCTTTCA GAGACGTCTG 2984
 AGGTCCTTTG CTCGGAGACT CCTCAGATCT GGAAACAGTG TGGTCCTTCC CATGCTGTAT 3044

20
 TCTGAAAAGA TCTCATATGG ATGTTGTGCT CTGAGCACCA CAGATGTGAT CTGCAGCTCG 3104
 TTTCTGAAAT GATGGAGTTC ATGGTGATCA GTGTGAGACT GGCCTCTCCC AGCAGGTTAA 3164
 AAATGGAGTT TTAAATTATA CTGTAGCTGA CAGTACTTCT GATTTTATAT TTATTTAGTC 3224

25
 TGAGTTGTAG AACTTTGCAA TCTAAGTTTA TTTTTTGTA CTAATAATT CATTTGGTGC 3284
 TGGTCTATTG ATTTTTGGGG GTAAACAATA TTATTCTTCA GAAGGGGACC TACTTCTTCA 3344
 TGGGAAGAAT TACTTTTATT CTCAAACCTAC AGAACAATGT GCTAAGCAGT GCTAAATTGT 3404

30
 TCTCATGAAG AAAACAGTCA CTGCATTTAT CTCTGTAGGC CTTTTTTCAG AGAGGCCTTG 3464
 TCTAGATTTT TGCCAGCTAG GCTACTGCAT GTCTTAGTGT CAGGCCTTAG GAAAGTGCCA 3524
 CGCTCTGCAC TAAAGATATC AGAGCTCTTG GTGTTACTTA GACAAGAGTA TGAGCAAGTC 3584

35
 GGACCTCTCA GAGTGTGGGA ACACAGTTTT GAAAGAAAAA CCATTTCTCT AAGCCAATTT 3644
 TCTTTAAAGA CATTTTAACT TATTTAGCTG AGTTCTAGAT TTTTCGGGTA AACTATCAAA 3704
 TCTGTATATG TTGTAATAAA GTGTCTTATG CTAGGAGTTT ATTCAAAGTG TTTAAGTAAT 3764

40
 AAAAGGACTC AAATTTACAC TGATAAAATA CTCTAGCTTG GGCCAGAGAA GACAGTGCTC 3824
 ATTAGCGTTG TCCAGGAAAC CCTGCTTGCT TGCCAAGCCT AATGAAGGGA AAGTCAGCTT 3884
 TCAGAGCCAA TGATGGAGGC CACATGAATG GCCCTGGAGC TGTGTGCCCT GTTCTGTGGC 3944

45
 CAGGAGCTTG GTGACTGAAT CATTTACGGG CTCCTTTGAT GGACCCATAA AAGCTCTTAG 4004
 CTTCTCAGG GGGTCAGCAG AGTTGTTGAA TCTTAATTTT TTTTAAATG TACCAGTTTT 4064

50

55

5
 GTATAAATAA TAATAAAGAG CTCCTTATTT TGTATTCTAT CTAATGCTTC GAGTTCAGTC 4124
 TTGGGAAGCT GACATCTCAT GTAGAAGATG GACTCTGAAA GACATTCCAA GAGTGCAGCG 4184
 10 GCATCATGGG AGCCTCTTAG TGATTGTGTG TCAGTATTAT TGTGGAAGAT TGACTTTGCT 4244
 TTTGTATGTG AAGTTTCAGA TTGCTCCTCT TGTGACTTTT TAGCCAGTAA CATTTTATTT 4304
 ACCTGAGCTT GTCATGGAAG TGGCAGTGAA AAGTATTGAG TATTCATGCT GGTGACTGTA 4364
 15 ACCAATGTCA TCTTGCTAAA AACTCATGTT TTGTACAATT ACTAAATTGT ATACATTTTG 4424
 TTATAGAATA CTTTTTCCAG TTGAGTAAAT TATGAAAGGA AGTTAACATT AACAGGTGTA 4484
 AGCGGTGGCT TTTTTAAAT GAAGGATTAA CCCTAAGCCC GAGACCCAGA AGCTAGCAAA 4544
 GTCTGGCAGA GTGGTAAACT GTCCTGCTGG GGCCATCCAA TCATCTCTCT CCATTACACT 4604
 20 TTCTAACTTT GCAGCATTGG TGCTGGCCAG TGTATTGTTT CATTGATCTT CCTTACGCTT 4664
 AGAGGGTTTG ATTGGTTCAG ATCTATAATC TCAGCCACAT TGTCTTGGTA TCAGCTGGAG 4724
 AGAGTTAAGA GGAAGGAAA ATAAAGTTCA GATAGCCAAA ACAC 4768

25

(2) INFORMATION FOR SEQ ID NO:4:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 887 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

30

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

35 Met Leu Ser Arg Leu Phe Arg Met His Gly Leu Phe Val Ala Ser His
 1 5 10 15
 Pro Trp Glu Val Ile Val Gly Thr Val Thr Leu Thr Ile Cys Met Met
 20 25 30
 40 Ser Met Asn Met Phe Thr Gly Asn Asn Lys Ile Cys Gly Trp Asn Tyr
 35 40 45
 Glu Cys Pro Lys Phe Glu Glu Asp Val Leu Ser Ser Asp Ile Ile Ile
 50 55 60
 45 Leu Thr Ile Thr Arg Cys Ile Ala Ile Leu Tyr Ile Tyr Phe Gln Phe
 65 70 75 80

50

55

5 Gln Asn Leu Arg Gln Leu Gly Ser Lys Tyr Ile Leu Gly Ile Ala Gly
 85 90 95
 Leu Phe Thr Ile Phe Ser Ser Phe Val Phe Ser Thr Val Val Ile His
 100 105 110
 10 Phe Leu Asp Lys Glu Leu Thr Gly Leu Asn Glu Ala Leu Pro Phe Phe
 115 120 125
 Leu Leu Leu Ile Asp Leu Ser Arg Ala Ser Ala Leu Ala Lys Phe Ala
 130 135 140
 15 Leu Ser Ser Asn Ser Gln Asp Glu Val Arg Glu Asn Ile Ala Arg Gly
 145 150 155 160
 Met Ala Ile Leu Gly Pro Thr Phe Thr Leu Asp Ala Leu Val Glu Cys
 165 170 175
 20 Leu Val Ile Gly Val Gly Thr Met Ser Gly Val Arg Gln Leu Glu Ile
 180 185 190
 Met Cys Cys Phe Gly Cys Met Ser Val Leu Ala Asn Tyr Phe Val Phe
 195 200 205
 25 Met Thr Phe Phe Pro Ala Cys Val Ser Leu Val Leu Glu Leu Ser Arg
 210 215 220
 Glu Ser Arg Glu Gly Arg Pro Ile Trp Gln Leu Ser His Phe Ala Arg
 225 230 235 240
 30 Val Leu Glu Glu Glu Glu Asn Lys Pro Asn Pro Val Thr Gln Arg Val
 245 250 255
 Lys Met Ile Met Ser Leu Gly Leu Val Leu Val His Ala His Ser Arg
 260 265 270
 35 Trp Ile Ala Asp Pro Ser Pro Gln Asn Ser Thr Thr Glu His Ser Lys
 275 280 285
 Val Ser Leu Gly Leu Asp Glu Asp Val Ser Lys Arg Ile Glu Pro Ser
 290 295 300
 40 Val Ser Leu Trp Gln Phe Tyr Leu Ser Lys Met Ile Ser Met Asp Ile
 305 310 315 320
 Glu Gln Val Val Thr Leu Ser Leu Ala Phe Leu Leu Ala Val Lys Tyr
 325 330 335
 45 Ile Phe Phe Glu Gln Ala Glu Thr Glu Ser Thr Leu Ser Leu Lys Asn
 340 345 350
 50
 55

5	Pro	Ile	Thr	Ser	Pro	Val	Val	Thr	Pro	Lys	Lys	Ala	Pro	Asp	Asn	Cys
			355					360					365			
	Cys	Arg	Arg	Glu	Pro	Leu	Leu	Val	Arg	Arg	Ser	Glu	Lys	Leu	Ser	Ser
	370						375					380				
10	Val	Glu	Glu	Glu	Pro	Gly	Val	Ser	Gln	Asp	Arg	Lys	Val	Glu	Val	Ile
	385					390					395					400
	Lys	Pro	Leu	Val	Val	Glu	Thr	Glu	Ser	Ala	Ser	Arg	Ala	Thr	Phe	Val
					405					410					415	
15	Leu	Gly	Ala	Ser	Gly	Thr	Ser	Pro	Pro	Val	Ala	Ala	Arg	Thr	Gln	Glu
				420					425					430		
	Leu	Glu	Ile	Glu	Leu	Pro	Ser	Glu	Pro	Arg	Pro	Asn	Glu	Glu	Cys	Leu
			435					440					445			
20	Gln	Ile	Leu	Glu	Ser	Ala	Glu	Lys	Gly	Ala	Lys	Phe	Leu	Ser	Asp	Ala
	450						455					460				
	Glu	Ile	Ile	Gln	Leu	Val	Asn	Ala	Lys	His	Ile	Pro	Ala	Tyr	Lys	Leu
	465					470					475					480
25	Glu	Thr	Leu	Met	Glu	Thr	His	Glu	Arg	Gly	Val	Ser	Ile	Arg	Arg	Gln
					485					490					495	
	Leu	Leu	Ser	Thr	Lys	Leu	Pro	Glu	Pro	Ser	Ser	Leu	Gln	Tyr	Leu	Pro
				500					505					510		
30	Tyr	Arg	Asp	Tyr	Asn	Tyr	Ser	Leu	Val	Met	Gly	Ala	Cys	Cys	Glu	Asn
			515					520					525			
	Val	Ile	Gly	Tyr	Met	Pro	Ile	Pro	Val	Gly	Val	Ala	Gly	Pro	Leu	Cys
	530						535					540				
35	Leu	Asp	Gly	Lys	Glu	Tyr	Gln	Val	Pro	Met	Ala	Thr	Thr	Glu	Gly	Cys
	545					550					555					560
	Leu	Val	Ala	Ser	Thr	Asn	Arg	Gly	Cys	Arg	Ala	Ile	Gly	Leu	Gly	Gly
					565					570					575	
40	Gly	Ala	Ser	Ser	Arg	Val	Leu	Ala	Asp	Gly	Met	Thr	Arg	Gly	Pro	Val
				580					585					590		
45	Val	Arg	Leu	Pro	Arg	Ala	Cys	Asp	Ser	Ala	Glu	Val	Lys	Ala	Trp	Leu
			595					600					605			
	Glu	Thr	Pro	Glu	Gly	Phe	Ala	Val	Ile	Lys	Asp	Ala	Phe	Asp	Ser	Thr
	610						615					620				

5 Ser Arg Phe Ala Arg Leu Gln Lys Leu His Val Thr Met Ala Gly Arg
 625 630 635 640
 Asn Leu Tyr Ile Arg Phe Gln Ser Lys Thr Gly Asp Ala Met Gly Met
 645 650 655
 10 Asn Met Ile Ser Lys Gly Thr Glu Lys Ala Leu Leu Lys Leu Gln Glu
 660 665 670
 Phe Phe Pro Glu Met Gln Ile Leu Ala Val Ser Gly Asn Tyr Cys Thr
 675 680 685
 15 Asp Lys Lys Pro Ala Ala Ile Asn Trp Ile Glu Gly Arg Gly Lys Thr
 690 695 700
 Val Val Cys Glu Ala Val Ile Pro Ala Lys Val Val Arg Glu Val Leu
 705 710 715 720
 20 Lys Thr Thr Thr Glu Ala Met Ile Asp Val Asn Ile Asn Lys Asn Leu
 725 730 735
 Val Gly Ser Ala Met Ala Gly Ser Ile Gly Gly Tyr Asn Ala His Ala
 740 745 750
 25 Ala Asn Ile Val Thr Ala Ile Tyr Ile Ala Cys Gly Gln Asp Ala Ala
 755 760 765
 Gln Asn Val Gly Ser Ser Asn Cys Ile Thr Leu Met Glu Ala Ser Gly
 770 775 780
 30 Pro Thr Asn Glu Asp Leu Tyr Ile Ser Cys Thr Met Pro Ser Ile Glu
 785 790 795 800
 Ile Gly Thr Val Gly Gly Gly Thr Asn Leu Leu Pro Gln Gln Ala Cys
 805 810 815
 35 Leu Gln Met Leu Gly Val Gln Gly Ala Cys Lys Asp Asn Pro Gly Glu
 820 825 830
 Asn Ala Arg Gln Leu Ala Arg Ile Val Cys Gly Thr Val Met Ala Gly
 835 840 845
 40 Glu Leu Ser Leu Met Ala Ala Leu Ala Ala Gly His Leu Val Arg Ser
 850 855 860
 His Met Val His Asn Arg Ser Lys Ile Asn Leu Gln Asp Leu Gln Gly
 865 870 875 880
 45 Thr Cys Thr Lys Lys Ser Ala
 885
 50
 55

5

(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 3348 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

10

(ii) MOLECULE TYPE: DNA (genomic)

15

- (ix) FEATURE:
 (A) NAME/KEY: CDS
 (B) LOCATION: 121..3255

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

20	GGAATATTTT GTACGAGCAA GTTATAGTAA GACACTTCAG TGAGAAATTA ATCTGACTTA	60
	CTTTTACTTA ATTGTGTTCT TTCCAAATTA GTTCAACAAG GTTCCACAT ACAACCTCAA	120
	ATG TCA CTT CCC TTA AAA ACG ATA GTA CAT TTG GTA AAG CCC TTT GCT	168
	Met Ser Leu Pro Leu Lys Thr Ile Val His Leu Val Lys Pro Phe Ala	
25	1 5 10 15	
	TGC ACT GCT AGG TTT AGT GCG AGA TAC CCA ATC CAC GTC ATT GTT GTT	216
	Cys Thr Ala Arg Phe Ser Ala Arg Tyr Pro Ile His Val Ile Val Val	
	20 25 30	
30	GCT GTT TTA TTG AGT GCC GCT GCT TAT CTA TCC GTG ACA CAA TCT TAC	264
	Ala Val Leu Leu Ser Ala Ala Ala Tyr Leu Ser Val Thr Gln Ser Tyr	
	35 40 45	
	CTT AAC GAA TGG AAG CTG GAC TCT AAT CAG TAT TCT ACA TAC TTA AGC	312
	Leu Asn Glu Trp Lys Leu Asp Ser Asn Gln Tyr Ser Thr Tyr Leu Ser	
35	50 55 60	
	ATA AAG CCG GAT GAG TTG TTT GAA AAA TGC ACA CAC TAC TAT AGG TCT	360
	Ile Lys Pro Asp Glu Leu Phe Glu Lys Cys Thr His Tyr Tyr Arg Ser	
	65 70 75 80	
40	CCT GTG TCT GAT ACA TGG AAG TTA CTC AGC TCT AAA GAA GCC GCC GAT	408
	Pro Val Ser Asp Thr Trp Lys Leu Leu Ser Ser Lys Glu Ala Ala Asp	
	85 90 95	
	ATT TAT ACC CCT TTT CAT TAT TAT TTG TCT ACC ATA AGT TTT CAA AGT	456
	Ile Tyr Thr Pro Phe His Tyr Tyr Leu Ser Thr Ile Ser Phe Gln Ser	
	100 105 110	

45

50

55

5

AAG GAC AAT TCA ACG ACT TTG CCT TCC CTT GAT GAC GTT ATT TAC AGT 504
Lys Asp Asn Ser Thr Thr Leu Pro Ser Leu Asp Asp Val Ile Tyr Ser
115 120 125

10

GTT GAC CAT ACC AGG TAC TTA TTA AGT GAA GAG CCA AAG ATA CCA ACT 552
Val Asp His Thr Arg Tyr Leu Leu Ser Glu Glu Pro Lys Ile Pro Thr
130 135 140

15

GAA CTA GTG TCT GAA AAC GGA ACG AAA TGG AGA TTG AGA AAC AAC AGC 600
Glu Leu Val Ser Glu Asn Gly Thr Lys Trp Arg Leu Arg Asn Asn Ser
145 150 155 160

AAT TTT ATT TTG GAC CTG CAT AAT ATT TAC CGA AAT ATG GTG AAG CAA 648
Asn Phe Ile Leu Asp Leu His Asn Ile Tyr Arg Asn Met Val Lys Gln
165 170 175

20

TTT TCT AAC AAA ACG AGC GAA TTT GAT CAG TTC GAT TTG TTT ATC ATC 696
Phe Ser Asn Lys Thr Ser Glu Phe Asp Gln Phe Asp Leu Phe Ile Ile
180 185 190

CTA GCT GCT TAC CTT ACT CTT TTT TAT ACT CTC TGT TGC CTG TTT AAT 744
Leu Ala Ala Tyr Leu Thr Leu Phe Tyr Thr Leu Cys Cys Leu Phe Asn
195 200 205

25

GAC ATG AGG AAA ATC GGA TCA AAG TTT TGG TTA AGC TTT TCT GCT CTT 792
Asp Met Arg Lys Ile Gly Ser Lys Phe Trp Leu Ser Phe Ser Ala Leu
210 215 220

30

TCA AAC TCT GCA TGC GCA TTA TAT TTA TCG CTG TAC ACA ACT CAC AGT 840
Ser Asn Ser Ala Cys Ala Leu Tyr Leu Ser Leu Tyr Thr Thr His Ser
225 230 235 240

TTA TTG AAG AAA CCG GCT TCC TTA TTA AGT TTG GTC ATT GGA CTA CCA 888
Leu Leu Lys Lys Pro Ala Ser Leu Leu Ser Leu Val Ile Gly Leu Pro
245 250 255

35

TTT ATC GTA GTA ATT ATT GGC TTT AAG CAT AAA GTT CGA CTT GCG GCA 936
Phe Ile Val Val Ile Ile Gly Phe Lys His Lys Val Arg Leu Ala Ala
260 265 270

TTC TCG CTA CAA AAA TTC CAC AGA ATT AGT ATT GAC AAG AAA ATA ACG 984
Phe Ser Leu Gln Lys Phe His Arg Ile Ser Ile Asp Lys Lys Ile Thr
275 280 285

40

GTA AGC AAC ATT ATT TAT GAG GCT ATG TTT CAA GAA GGT GCC TAC TTA 1032
Val Ser Asn Ile Ile Tyr Glu Ala Met Phe Gln Glu Gly Ala Tyr Leu
290 295 300

45

ATC CGC GAC TAC TTA TTT TAT ATT AGC TCC TTC ATT GGA TGT GCT ATT 1080
Ile Arg Asp Tyr Leu Phe Tyr Ile Ser Ser Phe Ile Gly Cys Ala Ile
305 310 315 320

50

55

5

	TAT	GCT	AGA	CAT	CTT	CCC	GGA	TTG	GTC	AAT	TTC	TGT	ATT	TTG	TCT	ACA	1128
	Tyr	Ala	Arg	His	Leu	Pro	Gly	Leu	Val	Asn	Phe	Cys	Ile	Leu	Ser	Thr	
					325					330					335		
10	TTT	ATG	CTA	GTT	TTC	GAC	TTG	CTT	TTG	TCT	GCT	ACT	TTT	TAT	TCT	GCC	1176
	Phe	Met	Leu	Val	Phe	Asp	Leu	Leu	Leu	Ser	Ala	Thr	Phe	Tyr	Ser	Ala	
				340					345					350			
	ATT	TTA	TCA	ATG	AAG	CTG	GAA	ATT	AAC	ATC	ATT	CAC	AGA	TCA	ACC	GTC	1224
	Ile	Leu	Ser	Met	Lys	Leu	Glu	Ile	Asn	Ile	Ile	His	Arg	Ser	Thr	Val	
15				355					360				365				
	ATC	AGA	CAG	ACT	TTG	GAA	GAG	GAC	GGA	GTT	GTC	CCA	ACT	ACA	GCA	GAT	1272
	Ile	Arg	Gln	Thr	Leu	Glu	Glu	Asp	Gly	Val	Val	Pro	Thr	Thr	Ala	Asp	
				370			375					380					
20	ATT	ATA	TAT	AAG	GAT	GAA	ACT	GCC	TCA	GAA	CCA	CAT	TTT	TTG	AGA	TCT	1320
	Ile	Ile	Tyr	Lys	Asp	Glu	Thr	Ala	Ser	Glu	Pro	His	Phe	Leu	Arg	Ser	
	385				390						395					400	
	AAC	GTG	GCT	ATC	ATT	CTG	GGA	AAA	GCA	TCA	GTT	ATT	GGT	CTT	TTG	CTT	1368
	Asn	Val	Ala	Ile	Ile	Leu	Gly	Lys	Ala	Ser	Val	Ile	Gly	Leu	Leu	Leu	
				405						410					415		
25	CTG	ATC	AAC	CTT	TAT	GTT	TTC	ACA	GAT	AAG	TTA	AAT	GCT	ACA	ATA	CTA	1416
	Leu	Ile	Asn	Leu	Tyr	Val	Phe	Thr	Asp	Lys	Leu	Asn	Ala	Thr	Ile	Leu	
				420					425					430			
	AAC	ACG	GTA	TAT	TTT	GAC	TCT	ACA	ATT	TAC	TCG	TTA	CCA	AAT	TTT	ATC	1464
30	Asn	Thr	Val	Tyr	Phe	Asp	Ser	Thr	Ile	Tyr	Ser	Leu	Pro	Asn	Phe	Ile	
			435					440					445				
	AAT	TAT	AAA	GAT	ATT	GGC	AAT	CTC	AGC	AAT	CAA	GTG	ATC	ATT	TCC	GTG	1512
	Asn	Tyr	Lys	Asp	Ile	Gly	Asn	Leu	Ser	Asn	Gln	Val	Ile	Ile	Ser	Val	
			450			455						460					
35	TTG	CCA	AAG	CAA	TAT	TAT	ACT	CCG	CTG	AAA	AAA	TAC	CAT	CAG	ATC	GAA	1560
	Leu	Pro	Lys	Gln	Tyr	Tyr	Thr	Pro	Leu	Lys	Lys	Tyr	His	Gln	Ile	Glu	
	465				470					475					480		
	GAT	TCT	GTT	CTA	CTT	ATC	ATT	GAT	TCC	GTT	AGC	AAT	GCT	ATT	CGG	GAC	1608
	Asp	Ser	Val	Leu	Leu	Ile	Ile	Asp	Ser	Val	Ser	Asn	Ala	Ile	Arg	Asp	
				485					490						495		
40	CAA	TTT	ATC	AGC	AAG	TTA	CTT	TTT	TTT	GCA	TTT	GCA	GTT	AGT	ATT	TCC	1656
	Gln	Phe	Ile	Ser	Lys	Leu	Leu	Phe	Phe	Ala	Phe	Ala	Val	Ser	Ile	Ser	
				500					505					510			
45	ATC	AAT	GTC	TAC	TTA	CTG	AAT	GCT	GCA	AAA	ATT	CAC	ACA	GGA	TAC	ATG	1704
	Ile	Asn	Val	Tyr	Leu	Leu	Asn	Ala	Ala	Lys	Ile	His	Thr	Gly	Tyr	Met	
			515					520					525				

50

55

5

AAC TTC CAA CCA CAA TCA AAT AAG ATC GAT GAT CTT GTT GTT CAG CAA 1752
 Asn Phe Gln Pro Gln Ser Asn Lys Ile Asp Asp Leu Val Val Gln Gln
 530 535 540

10

AAA TCG GCA ACG ATT GAG TTT TCA GAA ACT CGA AGT ATG CCT GCT TCT 1800
 Lys Ser Ala Thr Ile Glu Phe Ser Glu Thr Arg Ser Met Pro Ala Ser
 545 550 555 560

15

TCT GGC CTA GAA ACT CCA GTG ACC GCG AAA GAT ATA ATT ATC TCT GAA 1848
 Ser Gly Leu Glu Thr Pro Val Thr Ala Lys Asp Ile Ile Ile Ser Glu
 565 570 575

GAA ATC CAG AAT AAC GAA TGC GTC TAT GCT TTG AGT TCC CAG GAC GAG 1896
 Glu Ile Gln Asn Asn Glu Cys Val Tyr Ala Leu Ser Ser Gln Asp Glu
 580 585 590

20

CCT ATC CGT CCT TTA TCG AAT TTA GTG GAA CTT ATG GAG AAA GAA CAA 1944
 Pro Ile Arg Pro Leu Ser Asn Leu Val Glu Leu Met Glu Lys Glu Gln
 595 600 605

TTA AAG AAC ATG AAT AAT ACT GAG GTT TCG AAT CTT GTC GTC AAC GGT 1992
 Leu Lys Asn Met Asn Asn Thr Glu Val Ser Asn Leu Val Val Asn Gly
 610 615 620

25

AAA CTG CCA TTA TAT TCC TTA GAG AAA AAA TTA GAG GAC ACA ACT CGT 2040
 Lys Leu Pro Leu Tyr Ser Leu Glu Lys Lys Leu Glu Asp Thr Thr Arg
 625 630 635 640

30

GCG GTT TTA GTT AGG AGA AAG GCA CTT TCA ACT TTG GCT GAA TCG CCA 2088
 Ala Val Leu Val Arg Arg Lys Ala Leu Ser Thr Leu Ala Glu Ser Pro
 645 650 655

ATT TTA GTT TCC GAA AAA TTG CCC TTC AGA AAT TAT GAT TAT GAT CGC 2136
 Ile Leu Val Ser Glu Lys Leu Pro Phe Arg Asn Tyr Asp Tyr Asp Arg
 660 665 670

35

GTT TTT GGA GCT TGC TGT GAA AAT GTC ATC GGC TAT ATG CCA ATA CCA 2184
 Val Phe Gly Ala Cys Cys Glu Asn Val Ile Gly Tyr Met Pro Ile Pro
 675 680 685

40

GTT GGT GTA ATT GGT CCA TTA ATT ATT GAT GGA ACA TCT TAT CAC ATA 2232
 Val Gly Val Ile Gly Pro Leu Ile Ile Asp Gly Thr Ser Tyr His Ile
 690 695 700

CCA ATG GCA ACC ACG GAA GGT TGT TTA GTG GCT TCA GCT ATG CGT GGT 2280
 Pro Met Ala Thr Thr Glu Gly Cys Leu Val Ala Ser Ala Met Arg Gly
 705 710 715 720

45

TGC AAA GCC ATC AAT GCT GGT GGT GGT GCA ACA ACT GTT TTA ACC AAA 2328
 Cys Lys Ala Ile Asn Ala Gly Gly Gly Ala Thr Thr Val Leu Thr Lys
 725 730 735

50

55

5

	GAT	GGT	ATG	ACT	AGA	GGC	CCA	GTC	GTT	CGT	TTC	CCT	ACT	TTA	ATA	AGA	2376
	Asp	Gly	Met	Thr	Arg	Gly	Pro	Val	Val	Arg	Phe	Pro	Thr	Leu	Ile	Arg	
				740					745					750			
10	TCT	GGT	GCC	TGC	AAG	ATA	TGG	TTA	GAC	TCG	GAA	GAG	GGA	CAA	AAT	TCA	2424
	Ser	Gly	Ala	Cys	Lys	Ile	Trp	Leu	Asp	Ser	Glu	Glu	Gly	Gln	Asn	Ser	
			755					760					765				
15	ATT	AAA	AAA	GCT	TTT	AAT	TCT	ACA	TCA	AGG	TTT	GCA	CGT	TTG	CAA	CAT	2472
	Ile	Lys	Lys	Ala	Phe	Asn	Ser	Thr	Ser	Arg	Phe	Ala	Arg	Leu	Gln	His	
		770					775					780					
	ATT	CAA	ACC	TGT	CTA	GCA	GGC	GAT	TTG	CTT	TTT	ATG	AGA	TTT	CGG	ACA	2520
	Ile	Gln	Thr	Cys	Leu	Ala	Gly	Asp	Leu	Leu	Phe	Met	Arg	Phe	Arg	Thr	
						790					795					800	
20	ACT	ACC	GGT	GAC	GCA	ATG	GGT	ATG	AAC	ATG	ATA	TCG	AAA	GGT	GTC	GAA	2568
	Thr	Thr	Gly	Asp	Ala	Met	Gly	Met	Asn	Met	Ile	Ser	Lys	Gly	Val	Glu	
					805					810					815		
	TAC	TCT	TTG	AAA	CAA	ATG	GTA	GAA	GAA	TAT	GGT	TGG	GAA	GAT	ATG	GAA	2616
	Tyr	Ser	Leu	Lys	Gln	Met	Val	Glu	Glu	Tyr	Gly	Trp	Glu	Asp	Met	Glu	
25				820					825					830			
	GTT	GTC	TCC	GTA	TCT	GGT	AAC	TAT	TGT	ACT	GAT	AAG	AAA	CCT	GCC	GCA	2664
	Val	Val	Ser	Val	Ser	Gly	Asn	Tyr	Cys	Thr	Asp	Lys	Lys	Pro	Ala	Ala	
			835					840					845				
30	ATC	AAT	TGG	ATT	GAA	GGT	CGT	GGT	AAA	AGT	GTC	GTA	GCT	GAA	GCT	ACT	2712
	Ile	Asn	Trp	Ile	Glu	Gly	Arg	Gly	Lys	Ser	Val	Val	Ala	Glu	Ala	Thr	
		850					855					860					
	ATT	CCT	GGT	GAT	GTC	GTA	AAA	AGT	GTT	TTA	AAG	AGC	GAT	GTT	TCC	GCT	2760
	Ile	Pro	Gly	Asp	Val	Val	Lys	Ser	Val	Leu	Lys	Ser	Asp	Val	Ser	Ala	
		865				870					875					880	
35	TTA	GTT	GAA	TTA	AAT	ATA	TCC	AAG	AAC	TTG	GTT	GGA	TCC	GCA	ATG	GCT	2808
	Leu	Val	Glu	Leu	Asn	Ile	Ser	Lys	Asn	Leu	Val	Gly	Ser	Ala	Met	Ala	
					885					890					895		
	GGA	TCT	GTT	GGT	GGT	TTC	AAC	GCG	CAC	GCA	GCT	AAT	TTG	GTC	ACT	GCA	2856
40	Gly	Ser	Val	Gly	Gly	Phe	Asn	Ala	His	Ala	Ala	Asn	Leu	Val	Thr	Ala	
				900					905					910			
	CTT	TTC	TTG	GCA	TTA	GGC	CAA	GAT	CCT	GCG	CAG	AAC	GTC	GAA	AGT	TCC	2904
	Leu	Phe	Leu	Ala	Leu	Gly	Gln	Asp	Pro	Ala	Gln	Asn	Val	Glu	Ser	Ser	
			915					920					925				
45	AAC	TGT	ATA	ACT	TTG	ATG	AAG	GAA	GTT	GAT	GGT	GAT	TTA	AGG	ATC	TCT	2952
	Asn	Cys	Ile	Thr	Leu	Met	Lys	Glu	Val	Asp	Gly	Asp	Leu	Arg	Ile	Ser	
		930					935					940					

50

55

5

GTT TCC ATG CCA TCT ATT GAA GTT GGT ACG ATT GCC GGG GGT ACT GTT 3000
 Val Ser Met Pro Ser Ile Glu Val Gly Thr Ile Gly Gly Gly Thr Val 960
 945 950 955

10 CTG GAG CCT CAG GGC GCC ATG CTT GAT CTT CTC GGC GTT CGT GGT CCT 3048
 Leu Glu Pro Gln Gly Ala Met Leu Asp Leu Leu Gly Val Arg Gly Pro 975
 965 970

CAC CCC ACT GAA CCT GGA GCA AAT GCT AGG CAA TTA GCT AGA ATA ATC 3096
 His Pro Thr Glu Pro Gly Ala Asn Ala Arg Gln Leu Ala Arg Ile Ile 990
 980 985

15 GCG TGT GCT GTC TTG GCT GGT GAA CTG TCT CTG TGC TCC GCA CTT GCT 3144
 Ala Cys Ala Val Leu Ala Gly Glu Leu Ser Leu Cys Ser Ala Leu Ala 1005
 995 1000

20 GCC GGT CAC CTG GTA CAA AGC CAT ATG ACT CAC AAC CGT AAA ACA AAC 3192
 Ala Gly His Leu Val Gln Ser His Met Thr His Asn Arg Lys Thr Asn 1020
 1010 1015

AAA GCC AAT GAA CTG CCA CAA CCA AGT AAC AAA GGG CCC CCC TGT AAA 3240
 Lys Ala Asn Glu Leu Pro Gln Pro Ser Asn Lys Gly Pro Pro Cys Lys 1040
 1025 1030 1035

25 ACC TCA GCA TTA TTA TAACTCTTGT AGTTTACATG GTGATACTTT ATATCTTTGT 3295
 Thr Ser Ala Leu Leu 1045

ATTGTCTAGC TATTCTAAAT CATCTGCATG TAATAAGAAG TTGATCAAAA TGA 3348

30

(2) INFORMATION FOR SEQ ID NO:6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1045 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

35

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

40 Met Ser Leu Pro Leu Lys Thr Ile Val His Leu Val Lys Pro Phe Ala
 1 5 10 15

Cys Thr Ala Arg Phe Ser Ala Arg Tyr Pro Ile His Val Ile Val Val
 20 25 30

45 Ala Val Leu Leu Ser Ala Ala Ala Tyr Leu Ser Val Thr Gln Ser Tyr
 35 40 45

50

55

5 Leu Asn Glu Trp Lys Leu Asp Ser Asn Gln Tyr Ser Thr Tyr Leu Ser
 50 55 60
 Ile Lys Pro Asp Glu Leu Phe Glu Lys Cys Thr His Tyr Tyr Arg Ser
 65 70 75 80
 10 Pro Val Ser Asp Thr Trp Lys Leu Leu Ser Ser Lys Glu Ala Ala Asp
 85 90 95
 Ile Tyr Thr Pro Phe His Tyr Tyr Leu Ser Thr Ile Ser Phe Gln Ser
 100 105 110
 15 Lys Asp Asn Ser Thr Thr Leu Pro Ser Leu Asp Asp Val Ile Tyr Ser
 115 120 125
 Val Asp His Thr Arg Tyr Leu Leu Ser Glu Glu Pro Lys Ile Pro Thr
 130 135 140
 20 Glu Leu Val Ser Glu Asn Gly Thr Lys Trp Arg Leu Arg Asn Asn Ser
 145 150 155 160
 Asn Phe Ile Leu Asp Leu His Asn Ile Tyr Arg Asn Met Val Lys Gln
 165 170 175
 25 Phe Ser Asn Lys Thr Ser Glu Phe Asp Gln Phe Asp Leu Phe Ile Ile
 180 185 190
 Leu Ala Ala Tyr Leu Thr Leu Phe Tyr Thr Leu Cys Cys Leu Phe Asn
 195 200 205
 30 Asp Met Arg Lys Ile Gly Ser Lys Phe Trp Leu Ser Phe Ser Ala Leu
 210 215 220
 Ser Asn Ser Ala Cys Ala Leu Tyr Leu Ser Leu Tyr Thr Thr His Ser
 225 230 235 240
 35 Leu Leu Lys Lys Pro Ala Ser Leu Leu Ser Leu Val Ile Gly Leu Pro
 245 250 255
 Phe Ile Val Val Ile Ile Gly Phe Lys His Lys Val Arg Leu Ala Ala
 260 265 270
 40 Phe Ser Leu Gln Lys Phe His Arg Ile Ser Ile Asp Lys Lys Ile Thr
 275 280 285
 Val Ser Asn Ile Ile Tyr Glu Ala Met Phe Gln Glu Gly Ala Tyr Leu
 290 295 300
 45 Ile Arg Asp Tyr Leu Phe Tyr Ile Ser Ser Phe Ile Gly Cys Ala Ile
 305 310 315 320

50

55

5 Tyr Ala Arg His Leu Pro Gly Leu Val Asn Phe Cys Ile Leu Ser Thr
 325 330 335
 Phe Met Leu Val Phe Asp Leu Leu Leu Ser Ala Thr Phe Tyr Ser Ala
 340 345 350
 10 Ile Leu Ser Met Lys Leu Glu Ile Asn Ile Ile His Arg Ser Thr Val
 355 360 365
 Ile Arg Gln Thr Leu Glu Glu Asp Gly Val Val Pro Thr Thr Ala Asp
 370 375 380
 15 Ile Ile Tyr Lys Asp Glu Thr Ala Ser Glu Pro His Phe Leu Arg Ser
 385 390 395 400
 Asn Val Ala Ile Ile Leu Gly Lys Ala Ser Val Ile Gly Leu Leu Leu
 405 410 415
 20 Leu Ile Asn Leu Tyr Val Phe Thr Asp Lys Leu Asn Ala Thr Ile Leu
 420 425 430
 Asn Thr Val Tyr Phe Asp Ser Thr Ile Tyr Ser Leu Pro Asn Phe Ile
 435 440 445
 25 Asn Tyr Lys Asp Ile Gly Asn Leu Ser Asn Gln Val Ile Ile Ser Val
 450 455 460
 Leu Pro Lys Gln Tyr Tyr Thr Pro Leu Lys Lys Tyr His Gln Ile Glu
 465 470 475 480
 30 Asp Ser Val Leu Leu Ile Ile Asp Ser Val Ser Asn Ala Ile Arg Asp
 485 490 495
 Gln Phe Ile Ser Lys Leu Leu Phe Phe Ala Phe Ala Val Ser Ile Ser
 500 505 510
 35 Ile Asn Val Tyr Leu Leu Asn Ala Ala Lys Ile His Thr Gly Tyr Met
 515 520 525
 Asn Phe Gln Pro Gln Ser Asn Lys Ile Asp Asp Leu Val Val Gln Gln
 530 535 540
 40 Lys Ser Ala Thr Ile Glu Phe Ser Glu Thr Arg Ser Met Pro Ala Ser
 545 550 555 560
 45 Ser Gly Leu Glu Thr Pro Val Thr Ala Lys Asp Ile Ile Ile Ser Glu
 565 570 575
 Glu Ile Gln Asn Asn Glu Cys Val Tyr Ala Leu Ser Ser Gln Asp Glu
 580 585 590

50

55

5	Pro	Ile	Arg	Pro	Leu	Ser	Asn	Leu	Val	Glu	Leu	Met	Glu	Lys	Glu	Gln
			595					600					605			
	Leu	Lys	Asn	Met	Asn	Asn	Thr	Glu	Val	Ser	Asn	Leu	Val	Val	Asn	Gly
		610					615					620				
10	Lys	Leu	Pro	Leu	Tyr	Ser	Leu	Glu	Lys	Lys	Leu	Glu	Asp	Thr	Thr	Arg
	625					630					635					640
	Ala	Val	Leu	Val	Arg	Arg	Lys	Ala	Leu	Ser	Thr	Leu	Ala	Glu	Ser	Pro
					645					650					655	
15	Ile	Leu	Val	Ser	Glu	Lys	Leu	Pro	Phe	Arg	Asn	Tyr	Asp	Tyr	Asp	Arg
				660					665					670		
	Val	Phe	Gly	Ala	Cys	Cys	Glu	Asn	Val	Ile	Gly	Tyr	Met	Pro	Ile	Pro
			675					680					685			
20	Val	Gly	Val	Ile	Gly	Pro	Leu	Ile	Ile	Asp	Gly	Thr	Ser	Tyr	His	Ile
		690					695					700				
	Pro	Met	Ala	Thr	Thr	Glu	Gly	Cys	Leu	Val	Ala	Ser	Ala	Met	Arg	Gly
	705					710					715					720
25	Cys	Lys	Ala	Ile	Asn	Ala	Gly	Gly	Gly	Ala	Thr	Thr	Val	Leu	Thr	Lys
					725					730					735	
	Asp	Gly	Met	Thr	Arg	Gly	Pro	Val	Val	Arg	Phe	Pro	Thr	Leu	Ile	Arg
				740					745					750		
30	Ser	Gly	Ala	Cys	Lys	Ile	Trp	Leu	Asp	Ser	Glu	Glu	Gly	Gln	Asn	Ser
			755					760					765			
	Ile	Lys	Lys	Ala	Phe	Asn	Ser	Thr	Ser	Arg	Phe	Ala	Arg	Leu	Gln	His
		770					775					780				
35	Ile	Gln	Thr	Cys	Leu	Ala	Gly	Asp	Leu	Leu	Phe	Met	Arg	Phe	Arg	Thr
	785					790					795					800
40	Thr	Thr	Gly	Asp	Ala	Met	Gly	Met	Asn	Met	Ile	Ser	Lys	Gly	Val	Glu
					805					810					815	
	Tyr	Ser	Leu	Lys	Gln	Met	Val	Glu	Glu	Tyr	Gly	Trp	Glu	Asp	Met	Glu
				820					825					830		
45	Val	Val	Ser	Val	Ser	Gly	Asn	Tyr	Cys	Thr	Asp	Lys	Lys	Pro	Ala	Ala
			835					840					845			
	Ile	Asn	Trp	Ile	Glu	Gly	Arg	Gly	Lys	Ser	Val	Val	Ala	Glu	Ala	Thr
		850					855					860				

5 Ile Pro Gly Asp Val Val Lys Ser Val Leu Lys Ser Asp Val Ser Ala
 865 870 875 880
 Leu Val Glu Leu Asn Ile Ser Lys Asn Leu Val Gly Ser Ala Met Ala
 885 890 895
 10 Gly Ser Val Gly Gly Phe Asn Ala His Ala Ala Asn Leu Val Thr Ala
 900 905 910
 Leu Phe Leu Ala Leu Gly Gln Asp Pro Ala Gln Asn Val Glu Ser Ser
 915 920 925
 15 Asn Cys Ile Thr Leu Met Lys Glu Val Asp Gly Asp Leu Arg Ile Ser
 930 935 940
 Val Ser Met Pro Ser Ile Glu Val Gly Thr Ile Gly Gly Gly Thr Val
 945 950 955 960
 20 Leu Glu Pro Gln Gly Ala Met Leu Asp Leu Leu Gly Val Arg Gly Pro
 965 970 975
 His Pro Thr Glu Pro Gly Ala Asn Ala Arg Gln Leu Ala Arg Ile Ile
 980 985 990
 25 Ala Cys Ala Val Leu Ala Gly Glu Leu Ser Leu Cys Ser Ala Leu Ala
 995 1000 1005
 Ala Gly His Leu Val Gln Ser His Met Thr His Asn Arg Lys Thr Asn
 1010 1015 1020
 30 Lys Ala Asn Glu Leu Pro Gln Pro Ser Asn Lys Gly Pro Pro Cys Lys
 1025 1030 1035 1040
 Thr Ser Ala Leu Leu
 1045

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 22 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

GATCCGTCGA CGCATGCCTG CA

5

(2) INFORMATION FOR SEQ ID NO:8:

- 10 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 14 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

GGCATGCGTC GACG

14

20

(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
- 25 (ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

30 CCGGATCCGG

10

(2) INFORMATION FOR SEQ ID NO:10:

- 35 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 33 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)

40

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

AGCTTTCGCG AGCTCGAGAT CTAGATATCG ATG

33

(2) INFORMATION FOR SEQ ID NO:11:

- 45 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 32 base pairs

50

55

5

(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

10

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

15

AATTCATCGA TATCTAGATC TCGAGCTCGC GA

32

(2) INFORMATION FOR SEQ ID NO:12:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 24 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

20

(ii) MOLECULE TYPE: DNA (genomic)

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

TATCGAATTC AAGCTTGGTA CCGA

24

(2) INFORMATION FOR SEQ ID NO:13:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 24 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

30

(ii) MOLECULE TYPE: DNA (genomic)

35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

TATCGGTACC AAGCTTGAAT TCGA

24

40

(2) INFORMATION FOR SEQ ID NO:14:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 15 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

45

50

55

5

(ii) MOLECULE TYPE: DNA (genomic)

10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:
GATCCAGCTG TGTAC

15

(2) INFORMATION FOR SEQ ID NO:15:

15

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 17 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

20

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:
CCCCGGGATCG ATCACGT

17

25

(2) INFORMATION FOR SEQ ID NO:16

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 17 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

30

(ii) MOLECULE TYPE: DNA (genomic)

35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:
GATCGATCCC GGGACGT

17

(2) INFORMATION FOR SEQ ID NO:17:

40

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 30 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

45

50

55

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

10

ATAAAGACAT TGTTTTTAGA TCTGTTGTAA

30

(2) INFORMATION FOR SEQ ID NO:18:

15

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 32 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

20

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

GATTTATCTT CGTTTCCTGC AAGTTTTTGT TC

32

(2) INFORMATION FOR SEQ ID NO:19:

25

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 35 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

30

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

AGCTTCGAAG AACGAAGGAA GGAGCACAGA CTTAG

35

35

(2) INFORMATION FOR SEQ ID NO:20:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 34 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

40

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

45

ATTGGTATAT ATACGCATAT TGCGGCCGCG GTAC

34

50

55

5

(2) INFORMATION FOR SEQ ID NO:21:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 26 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

15

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

CGCGGCCGCA ATATGCGTAT ATATAC

26

20

(2) INFORMATION FOR SEQ ID NO:22:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 35 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

25

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

30

CAATCTAAGT CTGTGCTCCT TCCTTCGTTC TTCGA

35

(2) INFORMATION FOR SEQ ID NO:23:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 30 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

35

(ii) MOLECULE TYPE: DNA (genomic)

40

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

CTTTATGAGG GTAACATGAA TTCAAGAAGG

30

(2) INFORMATION FOR SEQ ID NO:24:

45

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 30 base pairs

50

55

(B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

GCCAAGTAGT TTTTACTCTT CAAGACAGAT AATTGCTGA CA

42

Claims

1. A method of increasing squalene, zymosterol, cholesta-7,24-dienol and cholesta-5,7,24-trienol accumulation in yeast comprising increasing the expression level of a structural gene encoding a polypeptide having HMG-CoA reductase activity in a mutant yeast having defects in the expression of zymosterol-24-methyltransferase and ergosta-5,7,24(28)-trienol-22-dehydrogenase.
2. The method according to claim 1 wherein said encoded polypeptide is an active, truncated HMG-CoA reductase enzyme.
3. The method according to claim 1 wherein said polypeptide is an active, truncated HMG-CoA reductase enzyme comprising the catalytic and at least a portion of the linker region but is free from the membrane binding region of S. cerevisiae HMG-CoA reductase #1.
4. The method according to claim 1 wherein said structural gene encodes an active, truncated HMG-CoA reductase enzyme comprising the catalytic and at least a portion of the linker region that is free from the membrane binding region of an HMG-CoA reductase enzyme.
5. The method according to claim 1 wherein the yeast is of the species S. cerevisiae.
6. The method according to claim 1 wherein squalene is accumulated relative to said zymosterol, cholesta-7,24-dienol and cholesta-5,7,24-trienol by culturing said yeast under conditions of restricted aeration.
7. The method according to claim 1 wherein the expression level is increased by increasing the copy number of a structural gene encoding a polypeptide having HMG-CoA reductase activity.
8. The method according to claim 7 wherein the copy number is increased by transforming said yeast with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes a polypeptide having HMG-CoA reductase activity, and a promoter suitable for driving the expression of said polypeptide in said yeast.
9. The method according to claim 8 wherein the promoter is selected from the group consisting of the GAL 1, GAL 10, GAL 1-10, PGK and ADH promoters.
10. The method according to claim 8 wherein the promoter and the exogenous DNA segment are integrated into the chromosomal DNA of said yeast.
11. A method of increasing squalene, ergosta-8,22-dienol, ergosta-7,22 dienol, ergosta-8-enol and ergosta-7-enol accumulation in yeast of the species S. cerevisiae comprising transforming a mutant S. cerevisiae having a defect in the expression of episterol-5-dehydrogenase with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes the catalytic region and at least a portion of the linker region but is free from the membrane binding region of an HMG-CoA reductase enzyme, and a promoter suitable for driving the expression of said reductase in said yeast.

- 5 12. A method of increasing squalene, zymosterol and cholesta-7,24-dienol accumulation in yeast of the species *S. cerevisiae* comprising transforming a mutant *S. cerevisiae* having a defect in the expression of zymosterol-24-methyltransferase and episterol-5-dehydrogenase with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes the catalytic region and at least a portion of the linker region but is free from the membrane binding region of an HMG-CoA reductase enzyme, and a promoter suitable for driving the expression of said reductase in said yeast.
- 10 13. A method of increasing squalene, zymosterol, ergosta-5,7,24(28)-trienol and ergosta-5,7-dienol accumulation in yeast of the species *S. cerevisiae* comprising transforming a mutant *S. cerevisiae* having a defect in the expression of ergosta-5,7,24(28)-trienol-22-dehydrogenase with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes the catalytic region and at least a portion of the linker region but is free from the membrane binding region of an HMG-CoA reductase enzyme, and a promoter suitable for driving the expression of said reductase in said yeast.
- 15 14. The method according to claim 11, 12 or 13 wherein the recombinant DNA molecule is selected from the group of plasmid vectors consisting of plasmids pSOC725ARC, pSOC106ARC, pARC300D, pARC306E, pARC300S, pARC300T and pARC304S.
- 20 15. A mutant *S. cerevisiae* having defects in the expression of zymosterol-24-methyltransferase and ergosta-5,7,24(28)-trienol-22-dehydrogenase enzymes, which mutant species is designated ATC0402mu.
- 25 16. A mutant of *S. cerevisiae* having single or double defects in the expression of enzymes that catalyze the conversion of squalene to ergosterol transformed with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes the catalytic region and at least a portion of the linker region but is free from the membrane binding region of an HMG-CoA reductase enzyme, and a promoter suitable for driving the expression of said reductase in said yeast.
- 30 17. The mutant according to claim 16 wherein the mutant is selected from the group consisting of mutants ATC0315rc, ATC1500, ATC1502, ATC1503, ATC1551, ATC2100, ATC2104, ACT2107, ACT2108, ATC2109 and ATC2401.
- 35 18. A recombinant DNA molecule selected from the group of plasmids designated plasmids pARC304S, pARC300S, pARC300T, pARC300D, pARC306E, pSOC106ARC and pSOC725ARC.
- 40
- 45
- 50
- 55

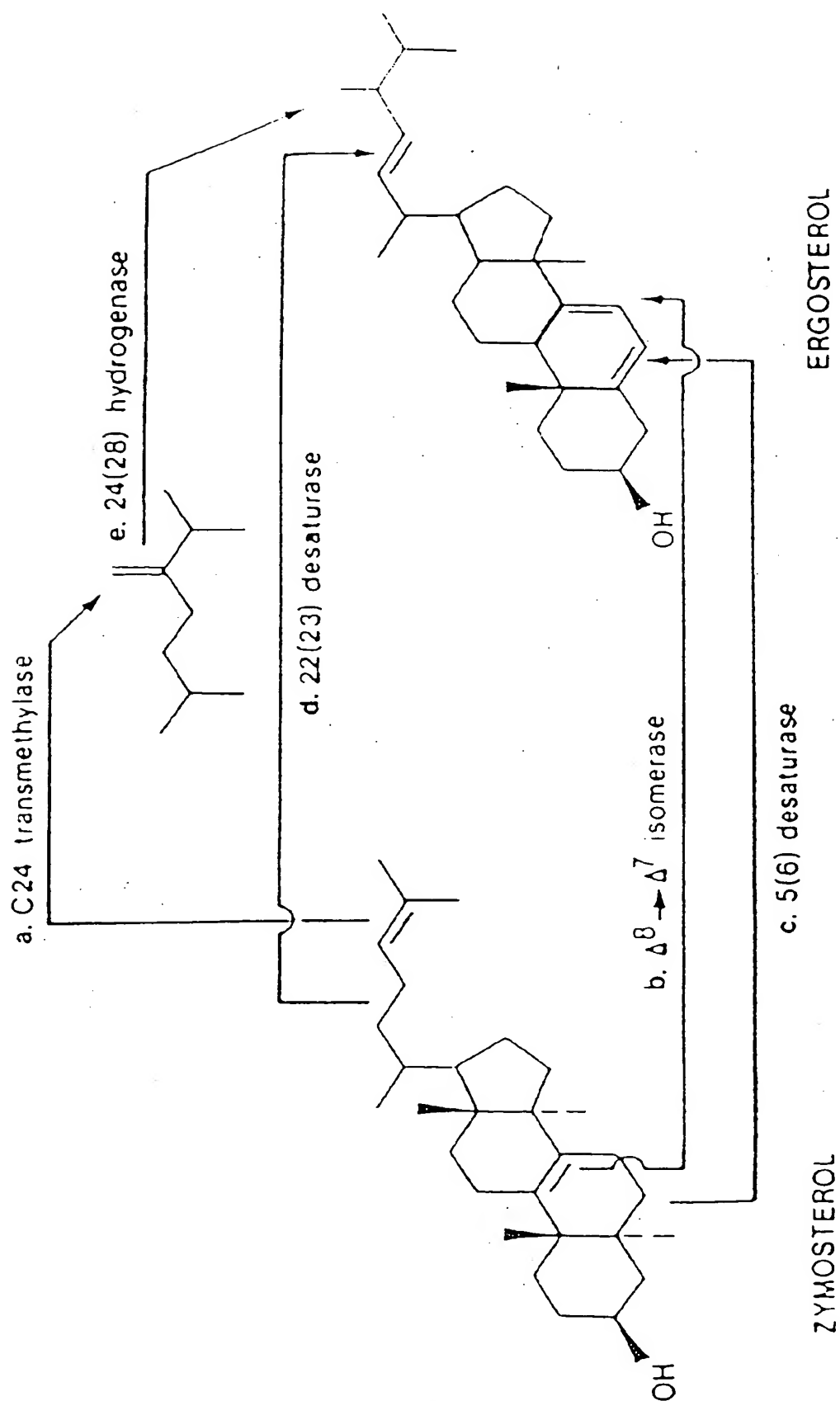


FIGURE 1

```

TTTATTAACT TATTTTTTTC TTCTTTCTAC CCAATTCTAG TCAGGAAAAG ACTAAGGGCT 60
GGAACATAGT GTATCATTGT CTAATTGTTG ATACAAAGTA GATAAATACA TAAACAAGC 120

ATG CCG CCG CTA TTC AAG GGA CTG AAA CAG ATG GCA AAG CCA ATT GCC 168
Met Pro Pro Leu Phe Lys Gly Leu Lys Gln Met Ala Lys Pro Ile Ala
1 5 10 15

TAT GTT TCA AGA TTT TCG GCG AAA CGA CCA ATT CAT ATA ATA CTT TTT 216
Tyr Val Ser Arg Phe Ser Ala Lys Arg Pro Ile His Ile Ile Leu Phe
20 25 30

TCT CTA ATC ATA TCC GCA TTC GCT TAT CTA TCC GTC ATT CAG TAT TAC 264
Ser Leu Ile Ile Ser Ala Phe Ala Tyr Leu Ser Val Ile Gln Tyr Tyr
35 40 45

TTC AAT GGT TGG CAA CTA GAT TCA AAT AGT GTT TTT GAA ACT GCT CCA 312
Phe Asn Gly Trp Gln Leu Asp Ser Asn Ser Val Phe Glu Thr Ala Pro
50 55 60

AAT AAA GAC TCC AAC ACT CTA TTT CAA GAA TGT TCC CAT TAC TAC AGA 360
Asn Lys Asp Ser Asn Thr Leu Phe Gln Glu Cys Ser His Tyr Tyr Arg
65 70 75 80

```

FIGURE 2-1

GAT TCC TCT CTA GAT GGT TGG GTA TCA ATC ACC GCG CAT GAA GCT AGT	408
Asp Ser Ser Leu Asp Gly Trp Val Ser Ile Thr Ala His Glu Ala Ser	85 90 95
GAG TTA CCA GCC CCA CAC CAT TAC TAT CTA TTA AAC CTG AAC TTC AAT	456
Glu Leu Pro Ala Pro His Tyr Tyr Leu Leu Asn Leu Asn Phe Asn	100 105 110
AGT CCT AAT GAA ACT GAC TCC ATT CCA GAA CTA GCT AAC ACG GTT TTT	504
Ser Pro Asn Glu Thr Asp Ser Ile Pro Glu Leu Ala Asn Thr Val Phe	115 120 125
GAG AAA GAT AAT ACA AAA TAT ATT CTG CAA GAA GAT CTC AGT GTT TCC	552
Glu Lys Asp Asn Thr Lys Tyr Ile Leu Gln Glu Asp Leu Ser Val Ser	130 135 140
AAA GAA ATT TCT TCT ACT GAT GGA GAT GGA ACG AAA TGG AGG TTA AGA AGT GAC	600
Lys Glu Ile Ser Ser Thr Asp Gly Thr Lys Trp Arg Leu Arg Ser Asp	145 150 155 160
AGA AAA AGT CTT TTC GAC GTA AAG ACG TTA GCA TAT TCT CTC TAC GAT	648
Arg Lys Ser Leu Phe Asp Val Lys Thr Leu Ala Tyr Ser Leu Tyr Asp	165 170 175

FIGURE 2-2

GTA TTT TCA GAA AAT GTA ACC CAA GCA GAC CCG TTT GAC GTC CTT ATT 696
 Val Phe Ser Glu Asn Val Thr Gln Ala Asp Pro Phe Asp Val Leu Ile 180 185 190

 ATG GTT ACT GCC TAC CTA ATG ATG TTC TAC ACC ATA TTC GGC CTC TTC 744
 Met Val Thr Ala Tyr Leu Met Met Phe Tyr Thr Ile Phe Gly Leu Phe 195 200 205

 AAT GAC ATG AGG AAG ACC GGG TCA AAT TTT TGG TTG AGC GCC TCT ACA 792
 Asn Asp Met Arg Lys Thr Gly Ser Asn Phe Trp Leu Ser Ala Ser Thr 210 215 220

 GTG GTC AAT TCT GCA TCA CTT TTC TTA GCA TTG TAT GTC ACC CAA 840
 Val Val Asn Ser Ala Ser Ser Leu Phe Leu Ala Leu Tyr Val Thr Gln 225 230 235 240

 TGT ATT CTA GGC AAA GAA GTT TCC GCA TTA ACT CTT TTT GAA GGT TTG 888
 Cys Ile Leu Gly Lys Glu Val Ser Ala Leu Thr Leu Phe Glu Gly Leu 245 250 255

 CCT TTC ATT GTA GTT GTT GGT TTC AAG CAC AAA ATC AAG ATT GCC 936
 Pro Phe Ile Val Val Val Val Gly Phe Lys His Lys Ile Lys Ile Ala 260 265 270

FIGURE 2-3

CAG TAT GCC CTG GAG AAA TTT GAA AGA GTC GGT TTA TCT AAA AGG ATT	984
Gln Tyr Ala Leu Glu Lys Phe Glu Arg Val Gly Leu Ser Lys Arg Ile	275 280 285
ACT ACC GAT GAA ATC GTT TTT GAA TCC GTG AGC GAA GAG GGT GGT CGT	1032
Thr Thr Asp Glu Ile Val Phe Glu Ser Val Ser Glu Gly Gly Arg	290 295 300
TTG ATT CAA GAC CAT TTG CTT TGT ATT TTT GCC TTT ATC GGA TGC TCT	1080
Leu Ile Gln Asp His Leu Leu Cys Ile Phe Ala Phe Ile Gly Cys Ser	305 310 315 320
ATG TAT GCT CAC CAA TTG AAG ACT TTG ACA AAC TTC TGC ATA TTA TCA	1128
Met Tyr Ala His Gln Leu Lys Thr Leu Thr Asn Phe Cys Ile Leu Ser	325 330 335
GCA TTT ATC CTA ATT TTT GAA TTG ATT TTA ACT CCT ACA TTT TAT TCT	1176
Ala Phe Ile Leu Ile Phe Glu Leu Ile Leu Thr Pro Thr Phe Tyr Ser	340 345 350
GCT ATC TTA GCG CTT AGA CTG GAA ATG AAT GTT ATC CAC AGA TCT ACT	1224
Ala Ile Leu Ala Leu Arg Leu Glu Met Asn Val Ile His Arg Ser Thr	355 360 365

FIGURE 2-4

ATT ATC AAG CAA ACA TTA GAA GAA GAC GGT GTT GTT CCA TCT ACA GCA	1272
Ile Ile Lys Gln Thr Leu Glu Glu Asp Gly Val Val Pro Ser Thr Ala	
370 375 380	
AGA ATC ATT TCT AAA GCA GAA AAG AAA TCC GTA TCT TCT TTC TTA AAT	1320
Arg Ile Ile Ser Lys Ala Glu Lys Lys Ser Val Ser Ser Phe Leu Asn	
385 390 395 400	
CTC AGT GTG GTT GTC ATT ATC ATG AAA CTC TCT GTC ATA CTG TTG TTT	1368
Leu Ser Val Val Val Ile Ile Met Lys Leu Ser Val Ile Leu Leu Phe	
405 410 415	
GTT TTC ATC AAC TTT TAT AAC TTT GGT GCA AAT TGG GTC AAT GAT GCC	1416
Val Phe Ile Asn Phe Tyr Asn Phe Gly Ala Asn Trp Val Asn Asp Ala	
420 425 430	
TTC AAT TCA TTG TAC TTC GAT AAG GAA CGT GGT TCT CTA CCA GAT TTT	1464
Phe Asn Ser Leu Tyr Phe Asp Lys Glu Arg Val Ser Leu Pro Asp Phe	
435 440 445	
ATT ACC TCG AAT GCC TCT GAA AAC TTT AAA GAG CAA GCT ATT GTT AGT	1512
Ile Thr Ser Asn Ala Ser Glu Asn Phe Lys Glu Gln Ala Ile Val Ser	
450 455 460	

FIGURE 2-5

GTC ACC CCA TTA TTA TAT TAC AAA CCC ATT AAG TCC TAC CAA CGC ATT	1560
Val Thr Pro Leu Leu Tyr Tyr TTA TTA TTA TTA TTA TTA TTA TTA TTA TTA	465
470	475
480	485
GAG GAT ATG GTT CTT CTA TTG CTT CGT AAT GTC AGT GTT GCC ATT CGT	1608
Glu Asp Met Val Leu Leu Leu Leu Arg Asn Val Ser Val Ala Ile Arg	490
485	495
GAT AGG TTC GTC AGT AAA TTA GTT CTT TCC GCC TTA GTA TGC AGT GCT	1656
Asp Arg Phe Val Ser Lys Leu Val Leu Ser Ala Leu Val Cys Ser Ala	500
505	510
GTC ATC AAT GTG TAT TTA TTG AAT GCT GCT AGA ATT CAT ACC AGT TAT	1704
Val Ile Asn Val Tyr Leu Leu Asn Ala Ala Arg Ile His Thr Ser Tyr	515
520	525
ACT GCA GAC CAA TTG GTG AAA ACT GAA GTC ACC AAG AAG TCT TTT ACT	1752
Thr Ala Asp Gln Leu Val Lys Thr Glu Val Thr Lys Lys Ser Phe Thr	530
535	540
GCT CCT GTA CAA AAG GCT TCT ACA CCA GTT TTA ACC AAT AAA ACA GTC	1800
Ala Pro Val Gln Lys Ala Ser Thr Pro Val Leu Thr Asn Lys Thr Val	545
550	555
560	565

FIGURE 2-6

ATT TCT GGA TCG AAA GTC AAA AGT TTA TCA TCT GCG CAA TCG AGC TCA	1848
Ile Ser Gly Ser Lys Val Lys Ser Leu Ser Ser Ala Gln Ser Ser Ser	575
	570
	565
TCA GGA CCT TCA TCA TCT AGT GAG GAA GAT GAT TCC CGC GAT ATT GAA	1896
Ser Gly Pro Ser Ser Ser Ser Glu Glu Asp Asp Ser Arg Asp Ile Glu	590
	585
	580
AGC TTG GAT AAG AAA ATA CGT CCT TTA GAA GAA TTA GCA TTA TTA	1944
Ser Leu Asp Lys Lys Ile Arg Pro Leu Glu Glu Ala Leu Leu	600
	595
	605
AGT AGT GGA AAT ACA AAA CAA TTG AAG AAC AAA GAG GTC GCT GCC TTG	1992
Ser Ser Gly Asn Thr Lys Lys Gln Leu Lys Asn Lys Glu Val Ala Ala Leu	610
	615
	620
GTT ATT CAC GGT AAG TTA CCT TTG TAC GCT TTG GAG AAA AAA TTA GGT	2040
Val Ile His Gly Lys Leu Pro Leu Tyr Ala Leu Glu Lys Lys Leu Gly	630
	635
	640
GAT ACT ACG AGA GCG GTT GCG GTA CGT AGG AAG GCT CTT TCA ATT TTG	2088
Asp Thr Thr Arg Ala Val Ala Val Arg Arg Lys Ala Leu Ser Ile Leu	645
	650
	655

FIGURE 2-7

GCA GAA GCT CCT GTA TTA GCA TCT GAT CGT TTA CCA TAT AAA AAT TAT	2136
Ala Glu Ala Pro Val Leu Ala Ser Asp Arg Leu Pro Tyr Lys Asn Tyr	
660 665 670	
GAC TAC GAC CGC GTA TTT GGC GCT TGT TGT GAA AAT GTT ATA GGT TAC	2184
Asp Tyr Asp Arg Val Phe Gly Ala Cys Cys Glu Asn Val Ile Gly Tyr	
675 680 685	
ATG CCT TTG CCC GTT GGT GTT ATA GGC CCC TTG GTT ATC GAT GGT ACA	2232
Met Pro Leu Pro Pro Val Gly Val Ile Ile Gly Pro Leu Val Ile Asp Gly Thr	
690 695 700	
TCT TAT CAT ATA CCA ATG GCA ACT ACA GAG GGT TGT TTG GTA GCT TCT	2280
Ser Tyr His Ile Pro Met Ala Thr Thr Glu Gly Cys Leu Val Ala Ser	
705 710 715 720	
GCC ATG CGT GGC TGT AAG GCA ATC AAT GCT GGC GGT GGT GCA ACA ACT	2328
Ala Met Arg Gly Cys Lys Ala Ile Asn Ala Gly Gly Gly Ala Thr Thr	
725 730 735	
GTT TTA ACT AAG GAT GGT ATG ACA AGA GGC CCA GTA GTC CGT TTC CCA	2376
Val Leu Thr Lys Asp Gly Met Thr Arg Gly Pro Val Val Arg Phe Pro	
740 745 750	

ACT TTG AAA AGA TCT GGT GCC TGT AAG ATA TGG TTA GAC TCA GAA GAG	2424
Thr Leu Lys Arg Ser Gly Ala Cys Lys Ile Trp Leu Asp Ser Glu Glu	755 760 765
GGA CAA AAC GCA ATT AAA AAA GCT TTT AAC TCT ACA TCA AGA TTT GCA	2472
Gly Gln Asn Ala Ile Lys Lys Ala Phe Asn Ser Thr Ser Arg Phe Ala	770 775 780
CGT CTG CAA CAT ATT CAA ACT TGT CTA GCA GGA GAT TTA CTC TTC ATG	2520
Arg Leu Gln His Ile Gln Thr Cys Leu Ala Gly Asp Leu Phe Met	785 790 795 800
AGA TTT AGA ACA ACT ACT GGT GAC GCA ATG GGT ATG AAT ATG ATT TCT	2568
Arg Phe Arg Thr Thr Thr Gly Asp Ala Met Gly Met Asn Met Ile Ser	805 810 815
AAA GGT GTC GAA TAC TCA TTA AAG CAA ATG GTA GAA GAG TAT GGC TGG	2616
Lys Gly Val Glu Tyr Ser Leu Lys Gln Met Val Glu Glu Tyr Gly Trp	820 825 830
GAA GAT ATG GAG GTT GTC TCC GTT TCT GGT AAC TAC TGT ACC GAC AAA	2664
Glu Asp Met Glu Val Val Ser Val Ser Gly Asn Tyr Cys Thr Asp Lys	835 840 845

FIGURE 2-9

AAA CCA GCT GCC ATC AAC TGG ATC GAA GGT CGT GGT AAG AGT GTC GTC	2712
Lys Pro Ala Ala Ile Asn Trp Ile Glu Gly Arg Gly Lys Ser Val Val	
850 855 860	
GCA GAA GCT ACT ATT CCT GGT GAT GTT GTC AGA AAA GTG TTA AAA AGT	2760
Ala Glu Ala Thr Ile Pro Gly Asp Val Val Arg Lys Val Leu Lys Ser	
865 870 875 880	
GAT GTT TCC GCA TTG GTT GAG TTG AAC ATT GCT AAG AAT TTG GTT GGA	2808
Asp Val Ser Ala Leu Val Glu Leu Asn Ile Ala Lys Asn Leu Val Gly	
885 890 895	
TCT GCA ATG GCT GGG TCT GTT GGT GGA TTT AAC GCA CAT GCA GCT AAT	2856
Ser Ala Met Ala Gly Ser Val Gly Gly Phe Asn Ala His Ala Ala Asn	
900 905 910	
TTA GTG ACA GCT GTT TTC TTG GCA TTA GGA CAA GAT CCT GCA CAA AAT	2904
Leu Val Thr Ala Val Phe Leu Ala Leu Gly Gln Asp Pro Ala Gln Asn	
915 920 925	
GTT GAA AGT TCC AAC TGT ATA ACA TTG ATG AAA GAA GTG GAC GGT GAT	2952
Val Glu Ser Ser Asn Cys Ile Thr Leu Met Lys Glu Val Asp Gly Asp	
930 935 940	

FIGURE 2-10

TTG AGA ATT TCC GTA TCC ATG CCA TCC ATC GAA GTA GGT ACC ATC GGT	3000
Leu Arg Ile Ser Val 950	
945	
GGT GGT ACT GTT CTA GAA CCA CCA GGT GCC ATG TTG GAC TTA TTA GGT	3048
Gly Gly Thr Val 965	
970	
GTA AGA GGC CCG CAT GCT ACC GCT CCT GGT ACC AAC GCA CGT CAA TTA	3096
Val Arg Gly Pro His Ala Thr Ala Pro Gly Thr Asn Ala Arg Gln Leu	
980	
985	
GCA AGA ATA GTT GCC TGT GCC GTC TTG GCA GGT GAA TTA TCC TTA TGT	3144
Ala Arg Ile Val Ala Cys Ala Val Leu Leu Ala Gly Glu Leu Ser Leu Cys	
995	
1000	
GCT GCC CTA GCA GCC GGC CAT TTG GTT CAA AGT CAT ATG ACC CAC AAC	3192
Ala Ala Leu Ala Ala Gly His Leu Val Gln Ser His Met Thr His Asn	
1010	
1015	
1020	
AGG AAA CCT GCT GAA CCA ACA AAA CCT AAC AAT TTG GAC GCC ACT GAT	3240
Arg Lys Pro Ala Glu Pro Thr Lys Pro Asn Asn Leu Asp Ala Thr Asp	
1025	
1030	
1035	
1040	

FIGURE 2-11

ATA AAT CGT TTG AAA GAT GGG TCC GTC ACC TGC ATT AAA TCC	3282
Ile Asn Arg Leu Lys Asp Gly Ser Val Thr Cys Ile Lys Ser	
1045	1050
TAAACTTAGT CATACGTCAT TGGTATTCTC TTGAAAAAGA AGCACAAACAG CACCATGTGT	3342
TACGTAAAT ATTACTT	3360

FIGURE 2-12

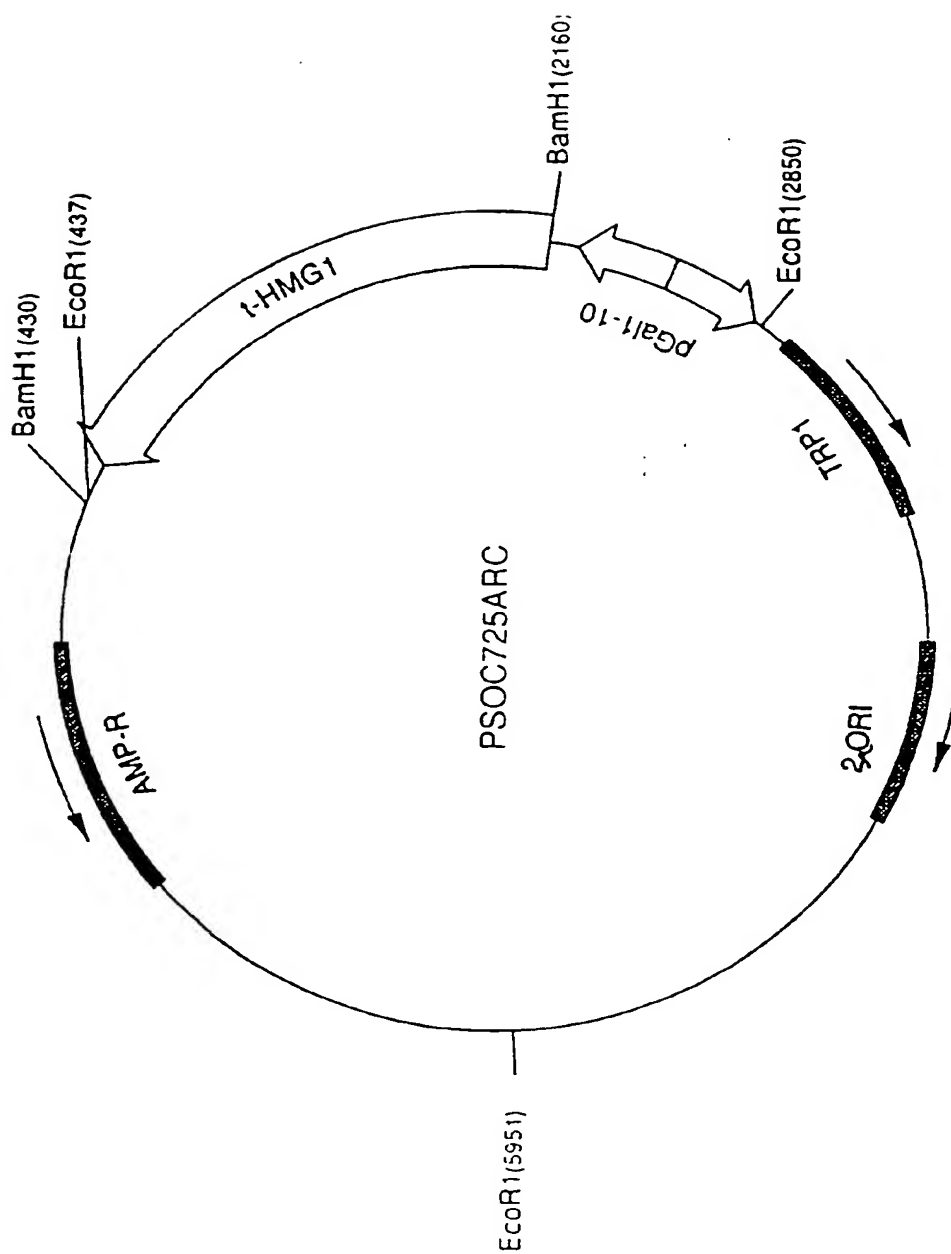


FIGURE 3

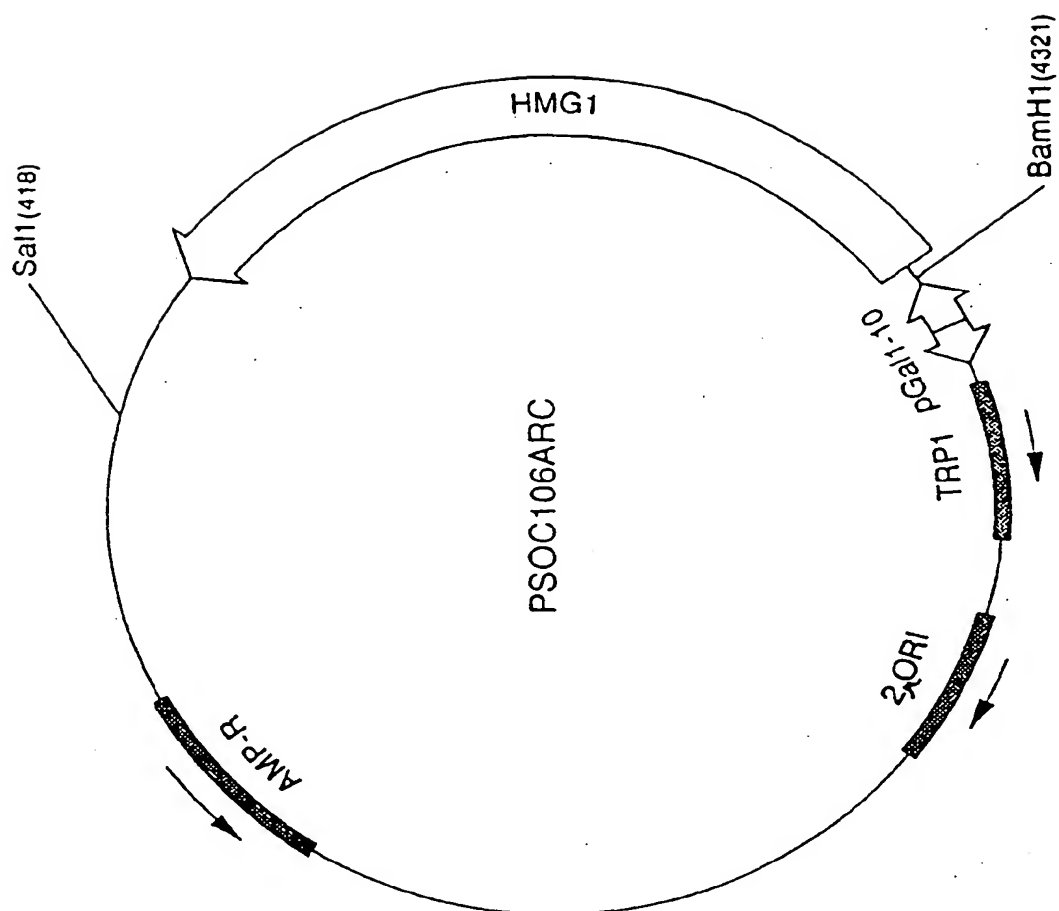


FIGURE 4

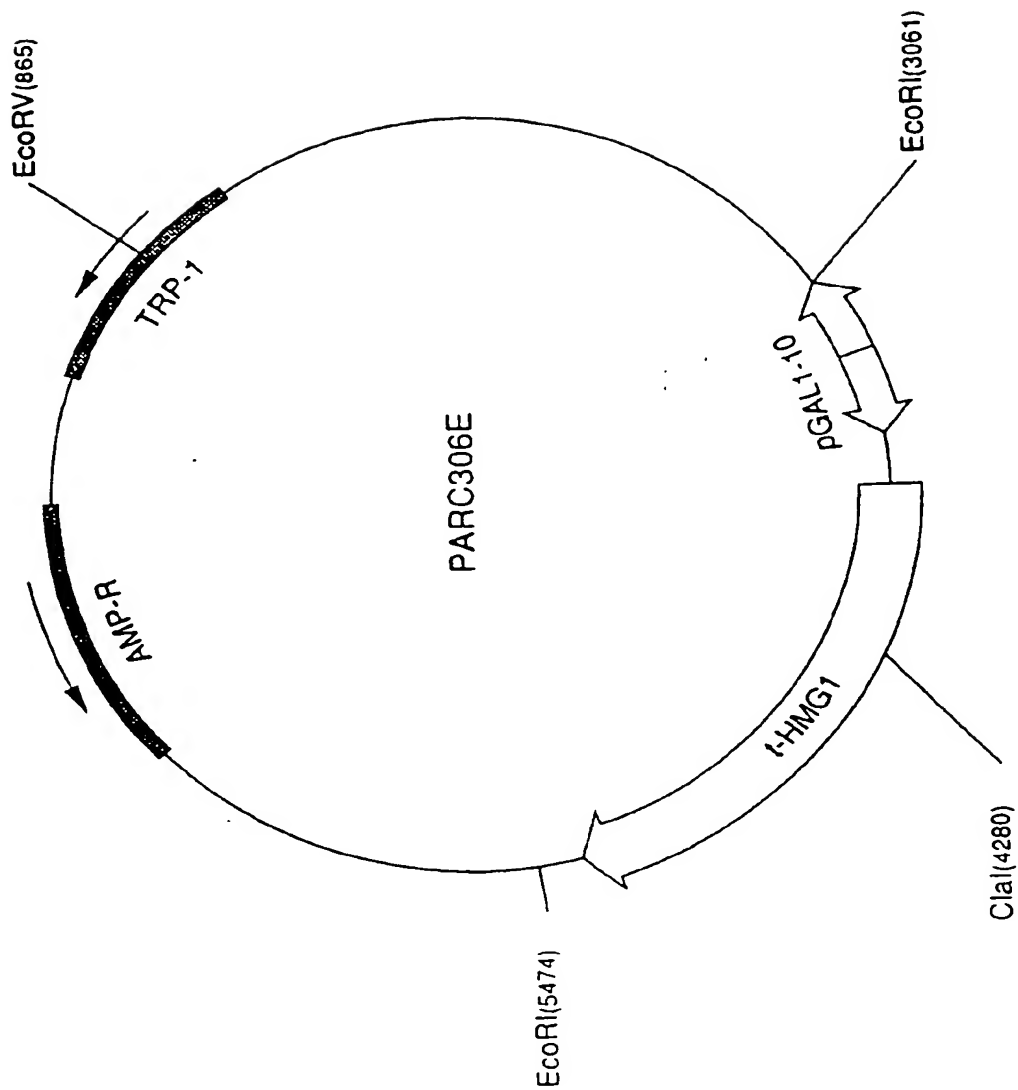


FIGURE 5

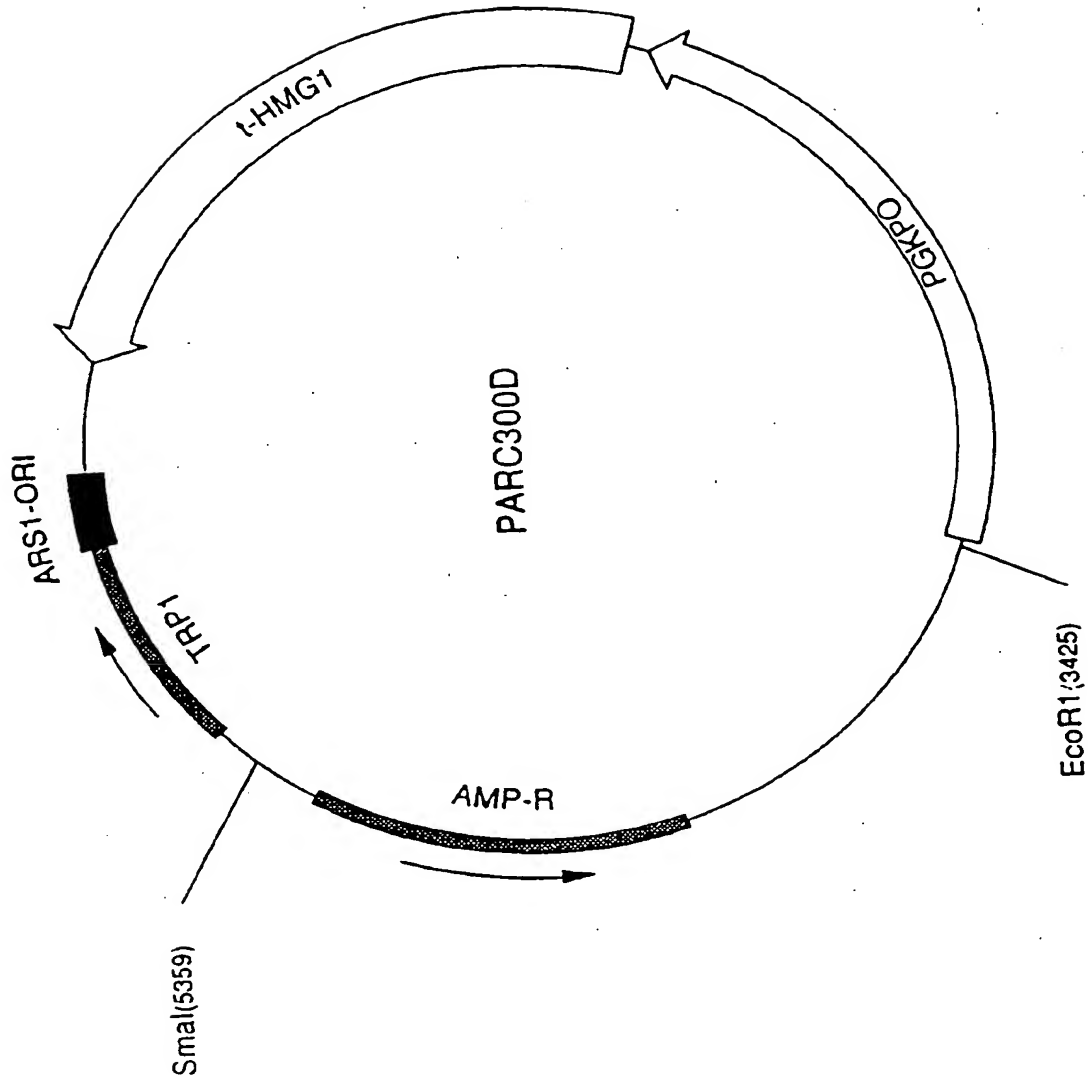


FIGURE 6

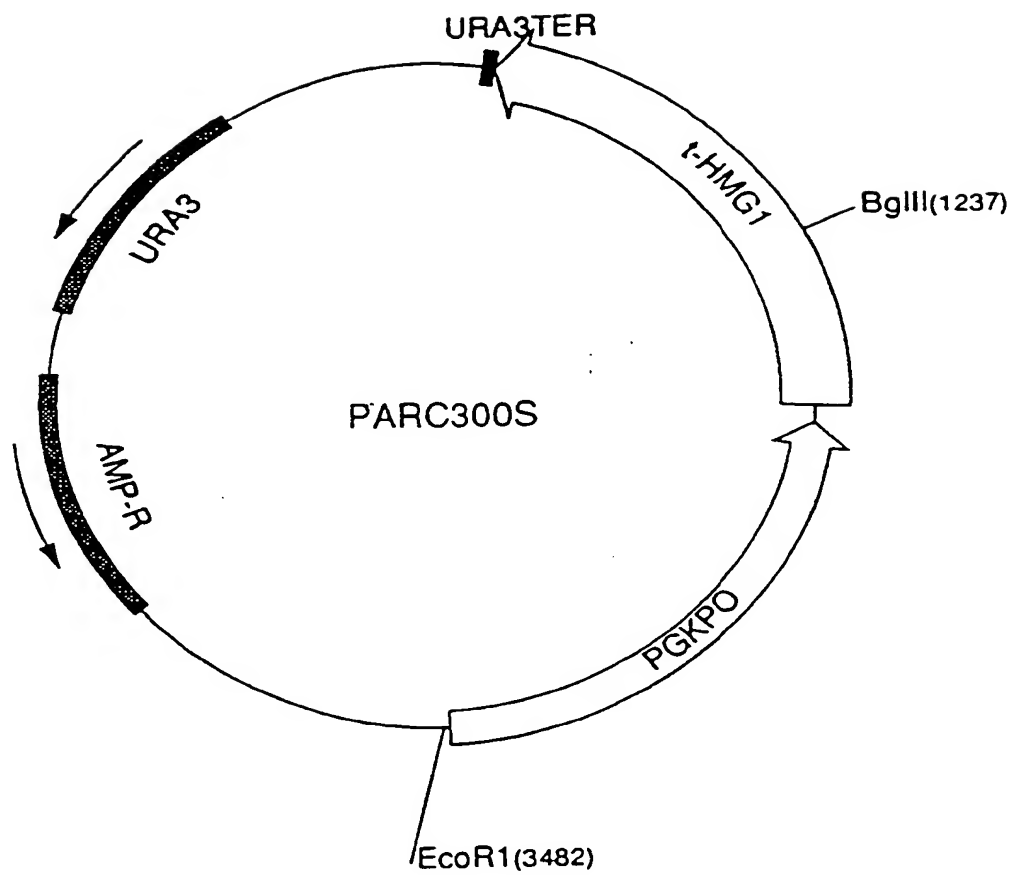


FIGURE 7

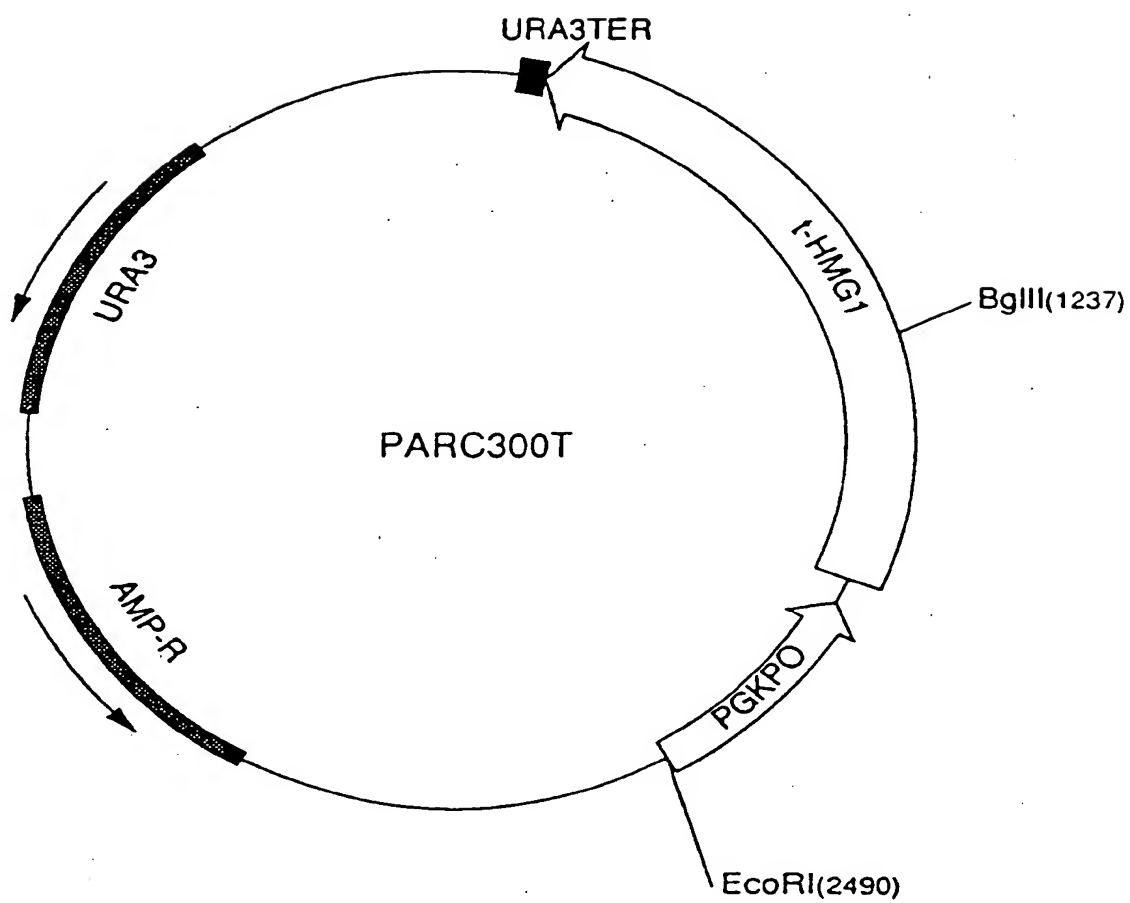


FIGURE 8

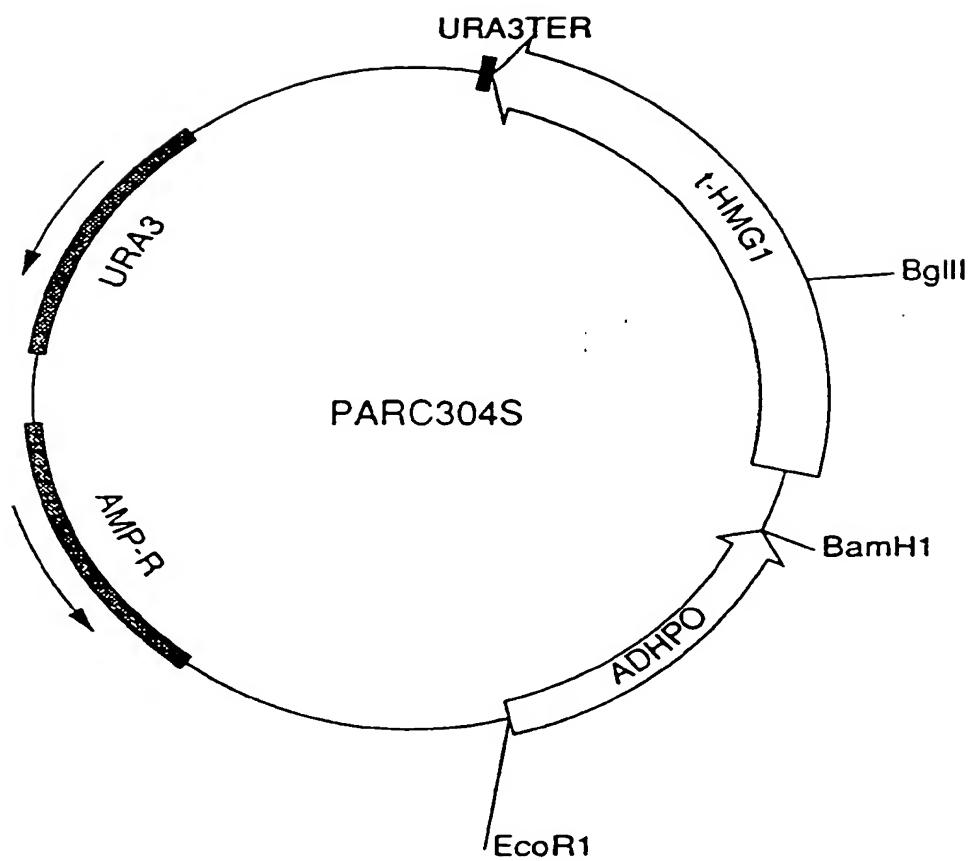


FIGURE 9



Europäisches Patentamt
European Patent Office
Office européen des brevets



Publication number : **0 486 290 A3**

(12)

EUROPEAN PATENT APPLICATION

(21) Application number : **91310494.9**

(22) Date of filing : **14.11.91**

(51) Int. Cl.⁵ : **C12N 15/81, C12N 15/53,
C12P 5/02, C12P 33/00,
C12N 1/19, C12N 1/18,
// (C12N1/18; C12R1:865),
(C12N1/19, C12R1:865)**

Declaration under Rule 28(4) EPC (expert solution)

(30) Priority : **15.11.90 US 613380
28.10.91 US 783861**

(43) Date of publication of application :
20.05.92 Bulletin 92/21

(84) Designated Contracting States :
BE CH DE DK ES FR GB IT LI NL

(88) Date of deferred publication of search report :
17.03.93 Bulletin 93/11

(71) Applicant : **AMOCO CORPORATION
200 East Randolph Drive
Chicago Illinois 60601 (US)**

(72) Inventor : **Saunders, Court A.
210 Holmes
Claredon Hills, IL 60514 (US)
Inventor : Wolf, Fred Richard
912 Muirhead Avenue
Naperville, IL 60565 (US)
Inventor : Mukharji, Indrani
2300 Central Park Avenue
Evanston, IL 60201 (US)**

(74) Representative : **Horton, Sophie Emma et al
Elkington and Fife Prospect House 8
Pembroke Road
Sevenoaks, Kent TN13 1XR (GB)**

(54) **A method and composition for increasing the accumulation of squalene and specific sterols in yeast.**

(57) A method of increasing the accumulation of squalene and specific sterols in yeast comprising increasing the expression level of a structural gene encoding a polypeptide having HMG-CoA reductase activity in a mutant yeast having single or double defects in the expression of sterol biosynthetic enzymes is provided. The expression level of a structural gene is preferably increased by transforming yeast with a recombinant DNA molecule comprising a vector operatively linked to an exogenous DNA segment that encodes a polypeptide having HMG-CoA reductase activity and a promoter that is suitable for driving the expression of the encoded polypeptide in the transformed yeast. The polypeptide having HMG-CoA reductase activity is preferably a truncated, active HMG-CoA reductase enzyme. Recombinant DNA molecules useful for transforming yeast and mutant yeast transformed with such recombinant DNA molecules are also disclosed.

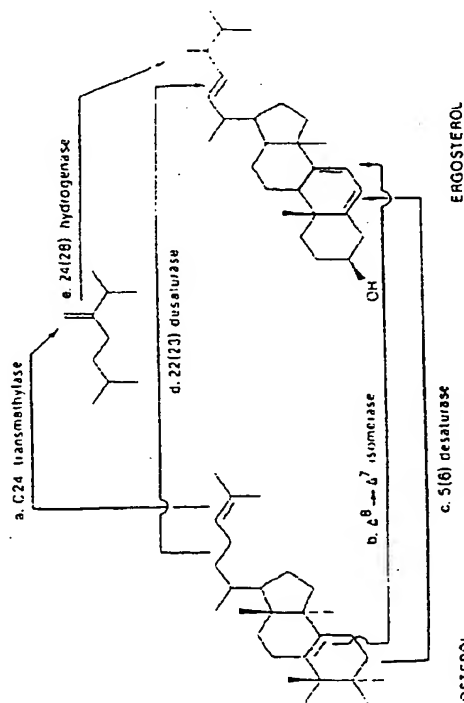


FIGURE 1

EP 0 486 290 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 91 31 0494

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
A	<p>BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS</p> <p>vol. 94, no. 3, 16 June 1980, DULUTH, MINNESOTA US</p> <p>pages 974 - 979</p> <p>JAMES F. DOWNING ET AL. 'The isolation of two mutants of <i>Saccharomyces cerevisiae</i> which demonstrate increased activity of 3-hydroxy-3-methylglutaryl coenzyme A reductase'</p> <p>* page 974, paragraph 1 - page 975, paragraph 1 *</p> <p>* page 977, paragraph 2 - page 978, paragraph 2 *</p> <p>-----</p>	1,5,6,15	<p>C12N15/81</p> <p>C12N15/53</p> <p>C12P5/02</p> <p>C12P33/00</p> <p>C12N1/19</p> <p>C12N1/18</p> <p>/(C12N1/18, C12R1:865)</p> <p>(C12N1/19, C12R1:865)</p>
			<p>TECHNICAL FIELDS SEARCHED (Int. Cl.5)</p>
			<p>C12N</p> <p>C12P</p> <p>C12R</p>
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 07 JANUARY 1993	Examiner MONTERO LOPEZ B.
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone</p> <p>Y : particularly relevant if combined with another document of the same category</p> <p>A : technological background</p> <p>O : non-written disclosure</p> <p>P : intermediate document</p> <p>T : theory or principle underlying the invention</p> <p>E : earlier patent document, but published on, or after the filing date</p> <p>D : document cited in the application</p> <p>L : document cited for other reasons</p> <p>*****</p> <p>& : member of the same patent family, corresponding document</p>			

EPO FORM 1503 03.82 (P0401)